

TRANSMITTAL LETTER TO THE UNITED STATES
 DESIGNATED/ELECTED OFFICE (DO/EO/US)
 CONCERNING A FILING UNDER 35 U.S.C. 371

Attorney's Docket Number

07385.0007

U.S. Application No.

09/529096

International Application. No.	International Filing Date	Priority Date Claimed
PCT/JP98/04671 ✓	October 15, 1998 ✓	October 17, 1997 ✓

Title of Invention:

AMIDE DERIVATIVES OR SALTS THEREOF ✓

Applicant(s) For DO/EO/US:

Tatsuya MARUYAMA, Takayuki SUZUKI, Kenichi ONDA, Masahiko HAYAKAWA, Hiroyuki MORITOMO,
 Tetsuya KIMIZUKA and Tetsuo MATSUI ✓

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US)
 the following items and other information:

1. [X] This is a FIRST submission of items concerning a filing under 35 U.S.C. 371.
2. [] This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. 371.
3. [] This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1).
4. [X] A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
5. [X] A copy of the International Application as filed (35 U.S.C. 371(c)(2))
 - a. [] is transmitted herewith (required only if not transmitted by the International Bureau).
 - b. [X] has been transmitted by the International Bureau.
 - c. [] is not required, as the application was filed in the United States Receiving Office (RO/US).
6. [X] A translation of the International Application into English (35 U.S.C. 371(c)(2)).
7. [X] Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3)).
 - a. [] are transmitted herewith (required only if not transmitted by the International Bureau).
 - b. [] have been transmitted by the International Bureau.
 - c. [] have not been made; however, the time limit for making such amendments has NOT expired.
 - d. [X] have not been made and will not be made.
8. [] A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. [X] An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. [] A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).

Items 11. to 16. below concern other document(s) or information included:

11. [] An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. [X] An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. [] A FIRST preliminary amendment.
14. [] A SECOND or SUBSEQUENT preliminary amendment.
15. [] A substitute specification.
16. [] A change of power of attorney and/or address letter.
16. [] Other items or information:
 - a. [] Verified Small Entity Statement.
 - b. [] Copy of Notification of Missing Requirements.

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422 Rec'd PCT/PTO 07 APR 2000

17. [X] The following fees are submitted:

CALCULATIONS

Basic National Fee (37 CFR 1.492(a)(1)-(5)):

Search Report has been prepared by the EPO or JPO.....\$840.00
 International preliminary examination fee paid to
 USPTO (37 CFR 1.482).....\$670.00
 No international preliminary examination fee paid to
 USPTO (37 CFR 1.482) but international search fee
 paid to USPTO (37 CFR 1.445(a)(2)).....\$690.00
 Neither international preliminary examination fee
 (37 CFR 1.482) nor international search fee
 (37 CFR 1.445(a)(2)) paid to USPTO.....\$970.00
 International preliminary examination fee paid to USPTO
 (37 CFR 1.482) and all claims satisfied provisions
 of PCT Article 33(1)-(4).....\$ 96.00

ENTER APPROPRIATE BASIC FEE AMOUNT = \$ 840.00

Surcharge of \$130.00 for furnishing the oath or declaration later than

[] 20 [] 30 months from the earliest claimed priority date

(37 CFR 1.492(e)).

Claims	Number Filed	Number Extra	Rate	
Total Claims	8 -20=		X \$18.00	\$
Independent Claims	3 - 3=		X \$78.00	\$
Multiple dependent claim(s) (if applicable)			+\$260.00	\$

TOTAL OF ABOVE CALCULATIONS = \$ 840.00

Reduction by 1/2 for filing by small entity, if applicable. Verified

Small Entity statement must also be filed. (Note 37 CFR 1.9, 1.27, 1.28)

SUBTOTAL = \$ 840.00

Processing fee of \$130.00 for furnishing the English translation later than [] 20 [] 30 months from the earliest claimed priority date

(37 CFR 1.492(f)).

TOTAL NATIONAL FEE = \$ 840.00

Fee for recording the enclosed assignment (37 CFR 1.21(h)). The

assignment must be accompanied by an appropriate cover sheet

(37 CFR 3.28, 3.31).

\$40.00 per property + \$ 40.00

TOTAL FEES ENCLOSED = \$ 880.00

Amount to be
 refunded \$
 charged \$

a. [X] A check in the amount of **\$880.00** to cover the above fees is enclosed.

b. [] Please charge my Deposit Account No. _____ in the amount of

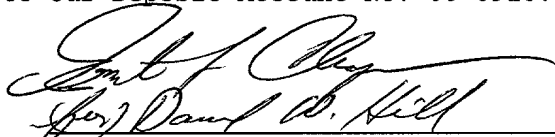
\$ _____

to cover the above fees. A duplicate copy of this sheet is enclosed.

c. [X] The Commissioner is hereby authorized to charge any additional fees
 which may be required, or credit any overpayment to Deposit Account
 No. 06-0916. A duplicate copy of this sheet is enclosed.

The Commissioner is hereby authorized to charge any other fees due under 37 C.F.R. §1.16
 or §1.17 during the pendency of this application to our Deposit Account No. 06-0916.

SEND ALL CORRESPONDENCE TO:
 Finnegan, Henderson, Farabow
 Garrett & Dunner, L.L.P.
 1300 I Street, N.W.
 Washington, D.C. 20005-3315
 DWH/FPD/rgm


 David W. Hill
 Reg. No. 28,220

Submitted: April 7, 2000

ERNEST F. CHAPMAN
 Reg. No. 25,961

Description

AMIDE DERIVATIVES OR SALTS THEREOF

Technical Field:

The present invention relates to pharmaceuticals and, more particularly, it relates to novel amide derivatives or salts thereof and also to therapeutic agents for diabetes mellitus containing them as effective components.

Background of the Invention:

Diabetes mellitus is a disease accompanied by continuous hyperglycemic state and is said to be resulted by action of many environmental factors and genetic factors. The main controlling factor for blood sugar is insulin, and it has been known that hyperglycemia is resulted by deficiency of insulin or by excess of factors which inhibit its action (such as genetic cause, lack of exercise, obesity and stress).

Diabetes mellitus is classified into two main types. One is insulin-dependent diabetes mellitus (IDDM) caused by a lowering of insulin-secreting function of pancreas due to autoimmune diseases, and another is non-insulin-dependent diabetes mellitus (NIDDM) caused by a lowering of insulin-secreting function of pancreas due to pancreatic fatigue accompanied by continuous high insulin secretion. 95% or more

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of diabetic patients in Japan are said to suffer from NIDDM, and an increase in the patients due to a change in daily life style is becoming a problem.

As to the therapy of diabetes mellitus, dietetic treatment, therapeutic exercise and remedy of obesity are mainly conducted in mild cases while, when the disease progresses, oral antidiabetic drugs (for example, insulin secretion promoters such as sulfonylurea compounds and insulin sensitivity potentiators which potentiate the sensitivity of insulin) are administered. In severe cases, an insulin preparation is administered. However, there has been a brisk demand for creation of the drugs whereby higher control for blood sugar is possible, and development of antidiabetic drugs having a new mechanism and having high usefulness has been demanded.

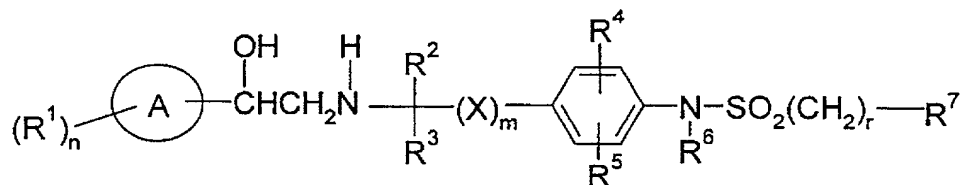
U.S. Patents 4,396,627 and 4,478,849 describe phenyl-ethanolamine derivatives and disclose that those compounds are useful as drugs for obesity and for hyperglycemia. Action of those compounds is reported to be due to a stimulating action to β_3 -receptors. Incidentally, it has been known that β -adrenaline receptors are classified into β_1 , β_2 and β_3 subtypes, that stimulation of β_1 -receptor causes an increase in heart rate, that stimulation of β_2 -receptor stimulates decomposition of glycogen in muscles, whereby synthesis of glycogen is inhibited, causing an action such as muscular tremor, and that stimulation of β_3 -receptor shows an anti-obesity and an

anti-hyperglycemia action (such as decrease in triglyceride, decrease in cholesterol and increase in HDL-cholesterol).

However, those β_3 -agonists also have actions caused by stimulation of β_1 - and β_2 -receptors such as increase in heart rate and muscular tremor, and they have a problem in terms of side effects.

Recently, it was ascertained that β -receptors have differences to species, and it has been reported that even compounds having been confirmed to have a β_3 -receptor selectivity in rodent animals such as rats show an action due to stimulating action to β_1 - and β_2 -receptors in human being. In view of the above, investigations for compounds having a stimulating action which is selective to β_3 -receptor in human being have been conducted recently using human cells or cells where human receptors are expressed. For example, WO 95/29159 describes substituted sulfonamide derivatives represented by the formula set forth below and discloses that due to their selective stimulating action to β_3 -receptors in human being, they are useful against obesity, hyperglycemia, etc. However, this patent does not specifically disclose an insulin secretion promoting action and an insulin sensitivity potentiating action of those compounds.

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(In the formula, the symbols should be referred to in the specification of this patent.)

As such, there has been still a demand for creation of therapeutic agents for diabetes mellitus of a new type which have a highly clinical usefulness.

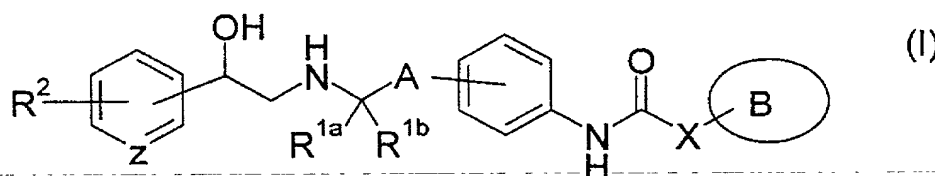
Disclosure of the Invention:

The present inventors have conducted an intensive investigation on compounds having both an insulin secretion promoting action and an insulin sensitivity potentiating action and found that novel amide derivatives show both a good insulin secretion promoting action and a good insulin sensitivity potentiating action and furthermore show a selective stimulating action to β_3 -receptors, leading to accomplishment of the present invention.

That is, the present invention relates to an amide derivative represented by the general formula (I) set forth below or a salt thereof that is useful for the therapy of diabetes mellitus, having both an insulin secretion promoting action and an insulin sensitivity potentiating action and further having anti-obesity and anti-hyperlipemia actions due

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to a selective stimulating action to β_3 -receptors. The present invention also relates to a pharmaceutical agent, particularly to a therapeutic agent for diabetes mellitus containing the amide derivative or the salt thereof as an effective ingredient.



(In the formula, each of the symbols means as follows:

ring B: a heteroaryl group which may be substituted and may be fused with a benzene ring;

X: a bond, lower alkylene or alkenylene which may be substituted with hydroxy or a lower alkyl group, carbonyl, or a group represented by -NH- (when X is a lower alkylene group which may be substituted with a lower alkyl group, the hydrogen atoms bonded to the carbon atom constituting the ring B may form a lower alkylene group together with the lower alkyl group so that a ring is formed);

A: lower alkylene or a group represented by -lower alkylene-O-;

R^{1a} , R^{1b} : they may be the same or different and each is a hydrogen atom or a lower alkyl group;

R²: a hydrogen atom or a halogen atom; and

Z: a nitrogen atom or a group represented by =CH-.)

The compound of the general formula (I) is further illustrated as follows.

In the definitions used in the general formula in this specification, the term "lower" means a linear or branched hydrocarbon chain having from 1 to 6 carbon atoms unless otherwise specified.

Specific examples of the "lower alkyl group" are methyl, ethyl, and linear or branched propyl, butyl, pentyl and hexyl, preferably an alkyl having from 1 to 4 carbon atoms, and particularly preferably methyl, ethyl, propyl and isopropyl.

Examples of the "lower alkylene group" is a divalent group obtained by removing an arbitrary hydrogen atom(s) from the above "lower alkyl group", preferably an alkylene group having from 1 to 4 carbon atoms, and particularly preferably methylene, ethylene, propylene and butylene. Examples of the "lower alkenylene group" are vinylene, propenylene, butenylene, pentenylene and hexenylene groups.

The "heteroaryl group which may be fused with a benzene ring" in the "heteroaryl group which may be substituted and may be fused with a benzene ring" means a ring group where a benzene ring is fused with a heteroaryl group as mentioned later or a non-fused heteroaryl group.

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Specific examples of the "ring group where the benzene ring is fused with a heteroaryl group" are fused-ring heteroaryl groups such as quinolyl, isoquinolyl, quinazolinyl, quinolidinyl, quinoxalinyl, cinnolinyl, benzimidazolyl, imidazopyridyl, benzofuranyl, benzoisoxazolyl, benzoxazolyl, benzothiazolyl, oxazolopyridyl, isothiazolopyridyl, benzothienyl, etc.; and oxo-added rings such as oxobenzofurayl, etc.

Examples of the "heteroaryl group" are monocyclic heteroaryl groups such as furyl, thienyl, pyrrolyl, imidazolyl, thiazolyl, pyrazolyl, isothiazolyl, isoxazolyl, pyridyl, pyrimidyl, pyridazinyl, pyrazinyl, thiadiazolyl, triazolyl, tetrazolyl, etc.; and bicyclic heteroaryl groups such as naphthylidinyll, pyridopyrimidinyl, etc.

The substituent in the "heteroaryl group which may be substituted and may be fused with a benzene ring" may be any group which can be usually substituted in this ring group. Preferred examples are a halogen atom and lower alkyl, lower alkenyl, lower alkynyl, hydroxy, sulfanyl, halogeno lower alkyl, lower alkyl-O-, lower alkyl-S-, lower alkyl-O-CO-, carboxy, sulfonyl, sulfinyl, lower alkyl-SO-, lower alkyl-SO₂-, lower alkyl-CO-, lower alkyl-CO-O-, carbamoyl, lower alkyl-NH-CO-, di-lower alkyl-N-CO-, nitro, cyano, amino, guanidino, lower alkyl-CO-NH-, lower alkyl-SO₂-NH-, lower alkyl-NH-, di-lower alkyl-N-, -O-lower alkylene-O-, etc. These substituents may further be substituted with a

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substituent such as an aryl group, a heteroaryl group, a halogen atom, hydroxy, sulfanyl, halogeno lower alkyl, lower alkyl-O-, lower alkyl-S-, lower alkyl-O-CO-, carboxy, sulfonyl, sulfinyl, lower alkyl-SO-, lower alkyl-SO₂-, lower alkyl-CO-, lower alkyl-CO-O-, carbamoyl, lower alkyl-NH-CO-, di-lower alkyl-N-CO-, nitro, cyano, amino, guanidino, lower alkyl-CO-NH-, lower alkyl-SO₂-NH-, lower alkyl-NH-, di-lower alkyl-N-, etc. These substituents such as an aryl group, a heteroaryl group, etc. may further be substituted with a halogen atom, etc.

The "lower alkenyl group" is a linear or branched alkenyl group having 2 to 6 carbon atoms, and its specific examples are vinyl, propenyl, butenyl, pentenyl and hexenyl groups.

The "lower alkynyl group" is a linear or branched alkynyl group having 2 to 6 carbon atoms, and its specific examples are ethynyl, propynyl, butynyl, pentynyl and hexynyl.

The "halogen atom" means a fluorine atom, a chlorine atom, a bromine atom or an iodine atom, and the "halogeno lower alkyl group" means a group where an arbitrary hydrogen atom or atoms in the above-mentioned alkyl group is/are substituted with a halogen atom or atoms.

The case when X is a bond means that a carbon atom of the -CO- group is directly bonded to the ring B.

The compound (I) of the present invention has at least one asymmetric carbon atom and therefore, there are optical

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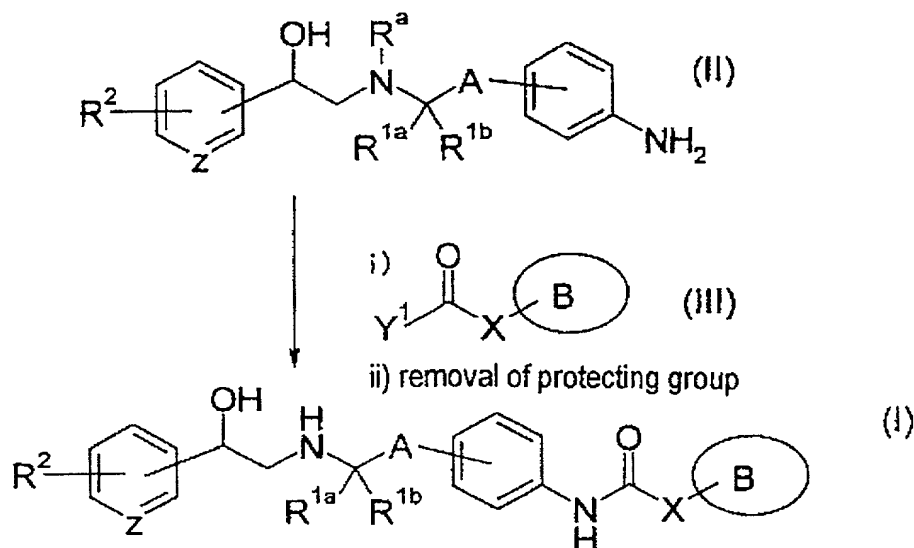
isomers such as (R)-compounds, (S)-compounds, etc., racemates, diastereomers, etc. The present invention includes all and each of isolated isomers and mixtures thereof. The present invention also includes hydrates, solvates (such as those with ethanol) and polymorphic substances of the compound (I).

The compound (I) of the present invention may form a salt with an acid. Examples of the salt are acid addition salts with mineral acids such as hydrochloric acid, hydrobromic acid, hydroiodic acid, sulfuric acid, nitric acid, phosphoric acid, etc.; and those with organic acids such as formic acid, acetic acid, propionic acid, oxalic acid, malonic acid, succinic acid, fumaric acid, maleic acid, lactic acid, malic acid, citric acid, tartaric acid, carbonic acid, picric acid, methanesulfonic acid, ethanesulfonic acid, glutamic acid, etc.

(Manufacturing Method)

The compound of the present invention or the salt thereof may be manufactured by application of various synthetic methods utilizing the characteristics of its fundamental skeleton or type of the substituent. Representative manufacturing methods are illustrated as hereunder.

First Manufacturing Method:



(In the formulae, R^{1a} , R^{1b} , R^2 , A , B , X and Z have the same meanings as defined already; R^a is a protective group for amino; and Y^1 is a leaving group, and more specifically hydroxy, lower alkoxy or halide.)

In this method, the compound (II) and the compound (III) are subjected to amidation, and the protective group is then removed therefrom to synthesize the compound (I) of the present invention.

The amidation in this manufacturing method can be conducted by customary manners.

The solvent may vary depending upon Y^1 of the compound (III) and mostly, an inert solvent or an alcoholic solvent (such as isopropanol, etc.) may be applied.

When Y^1 is a hydroxy group, a method where the reaction is conducted in the above-mentioned solvent in the presence of a condensing agent may be applied. Examples of the condensing agent are N,N'-dicyclohexylcarbodiimide (DCC), 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide (EDCI), 1,1'-carbonyldiimidazole (CDI), diphenylphosphoryl azide (DPPA), diethylphosphoryl cyanide (DEPC), etc.

When Y^1 is lower alkoxy, a method where the reaction is conducted under heating or refluxing as it is or in the above-mentioned inert solvent may be applied.

When Y^1 is halide, a method where the reaction is conducted in the above-mentioned inert solvent in the presence of a base may be applied.

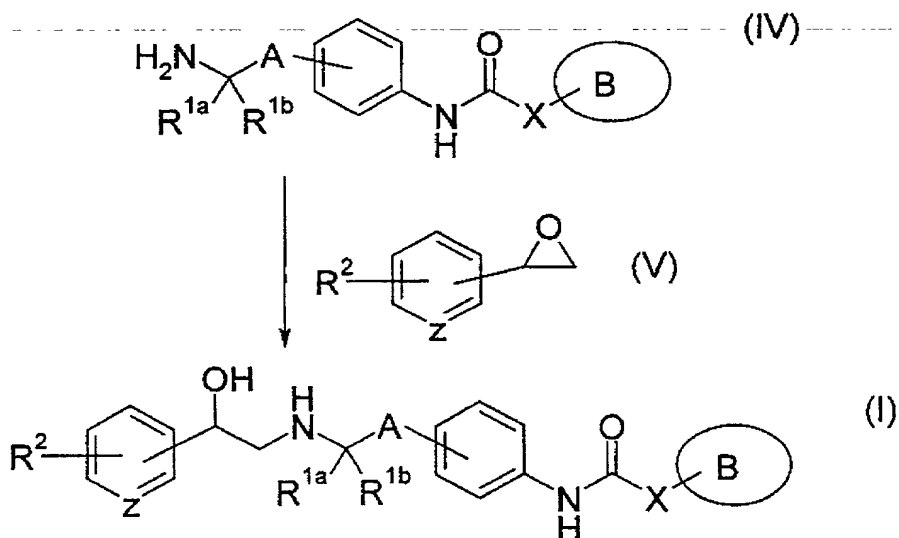
Examples of the inert solvent are dimethylformamide (DMF), dimethylacetamide, tetrachloroethane, dichloromethane, dichloroethane, chloroform, carbon tetrachloride, tetrahydrofuran, dioxane, dimethoxyethane, ethyl acetate, benzene, toluene, xylene, acetonitrile, dimethyl sulfoxide, etc., and mixed solvents thereof, and they may be appropriately selected depending upon each reaction condition. Examples of the base are inorganic bases such as sodium hydroxide, potassium hydroxide, sodium carbonate, potassium carbonate, etc.; and organic bases such as N-methylmorpholine, triethylamine, diisopropylethylamine, pyridine, etc.

The protective group of the amino represented by R^a means a protective group which is commonly used for amino by those skilled in the art, and its representative examples are acyl such as formyl, acetyl, propionyl, methoxyacetyl, methoxypropionyl, benzoyl, thienylacetyl, thiazolylacetyl, tetrazolylacetyl, thiazolylglyoxyloyl, thienylglyoxyloyl, etc.; lower alkoxy-carbonyl such as methoxycarbonyl, ethoxycarbonyl, tert-butoxycarbonyl, etc.; aralkyloxy-carbonyl such as benzyloxycarbonyl, p-nitrobenzyloxycarbonyl, etc.; lower alkanesulfonyl such as methanesulfonyl, ethanesulfonyl, etc.; aralkyl such as benzyl, p-nitrobenzyl, benzhydryl, trityl, etc.; tri-(lower alkyl)silyl such as trimethylsilyl, etc.; and the like.

Removal of the protective group in this manufacturing method may be conducted by customary manners. For example, the protective group for amino represented by R^a may be easily removed, for example, by i) a method where in case that the protective group is benzhydryl, p-methoxybenzyl, trityl, tert-butoxycarbonyl, formyl, etc., treatment with an acid such as formic acid, trifluoroacetic acid, a trifluoroacetic acid-anisole mixed solution, a hydrobromic acid-acetic acid mixed solution, a hydrochloric acid-dioxane mixed solution, etc. is conducted; ii) a method where in case that the protective group is benzyl, p-nitrobenzyl, benzhydryl, trityl, etc., a catalytic reduction method using palladium-carbon or

palladium hydroxide-carbon is conducted; and iii) a method where in case that the protective group is a tri-(lower alkyl)silyl or the like, treatment with water, fluoride anion (e.g., tetra-n-butylammonium fluoride, sodium fluoride, potassium fluoride, hydrofluoric acid), etc. is conducted.

Second Manufacturing Method:



(In the formulae, R^{1a} , R^{1b} , R^2 , A , B , X and Z have the same meanings as defined already.)

In this manufacturing method, the compound (IV) is reacted with the compound (V) to give the compound (I) of the present invention.

The amine compound (IV) and the compound (V) are reacted under heating or refluxing for 1 to 24 hours as they are or in an inert solvent, to give the compound (I) of the present invention.

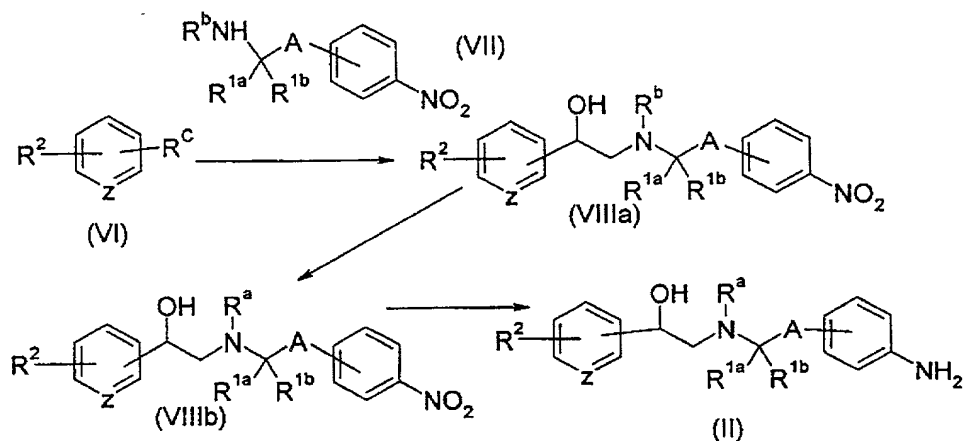
Examples of the inert solvent are acetonitrile, tetrahydrofuran, 2-butanone, dimethyl sulfoxide and N-methylpyrrolidone. In the reaction, a base such as sodium bicarbonate, potassium carbonate or diisopropylethylamine may be added to the reaction mixture.

Incidentally, in the above manufacturing methods, it is possible to purify the resulting substance by removing undesired by-products by means of recrystallization, pulverization, preparative thin layer chromatography, silica gel flash chromatography (as described in W. C. Still, et al., *J. Org. Chem.*, **43**, 2923 (1978)), medium-pressure liquid chromatography and HPLC. The compound produced through HPLC can be isolated as a corresponding salt.

The starting material used in the above-mentioned manufacturing methods may be easily manufactured by the methods which are known to those skilled in the art. One of the representative methods is shown as hereunder.

(Manufacturing Method for the Starting Compound (II))

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(In the formulae, R^{1a} , R^{1b} , R^2 , R^a , A and Z have the same meanings as defined already; R^b is a hydrogen atom or an aralkyl-based protective group for amino; and R^C is epoxy, 2-haloacetyl or 1-carboxymethan-1-ol.)

This manufacturing method is composed of from step (a) to step (c) in which the step (a) is a step where the compound (VI) is reacted with the compound (VII), followed by reduction reaction to give the compound (VIIIa) depending upon the type of R^C ; the step (b) is a step where protection is conducted when R^b of the compound (VIIIa) is a hydrogen atom; and the step (c) is a step where nitro is reduced to amino to give the compound (II).

Examples of the aralkyl-based protective group for amino used in this manufacturing method are benzyl, p-nitrobenzyl, benzhydryl, etc.

Step (a):

Illustration is made for the following three cases.

1) When R^c is epoxy, the compound (VI) may be reacted with the compound (VII) by the same manner as in the above-mentioned second manufacturing method. Reaction conditions such as reaction temperature, solvent, etc. are the same as well.

2) When R^c is 2-haloacetyl, the compound (VI) is reacted with the compound (VII) in the presence of a base, followed by reduction reaction to prepare the compound (VIIIa). The base is the same as that mentioned in the first manufacturing method. The reduction reaction may be conducted in the above-mentioned inert solvent or in a solvent of an alcohol type with stirring in the presence of a reducing agent. Examples of the reducing agent are sodium borohydride, sodium cyanoborohydride, lithium aluminum hydride, borane, etc.

3) When R^c is 1-carboxymethan-1-ol, the compound (VI) is reacted with the compound (VII) in the presence of a condensing agent, followed by reduction reaction in the same manner as in 2) to prepare the compound (VIIIa). The condensing agent is the same as that mentioned in the first manufacturing method.

Step (b):

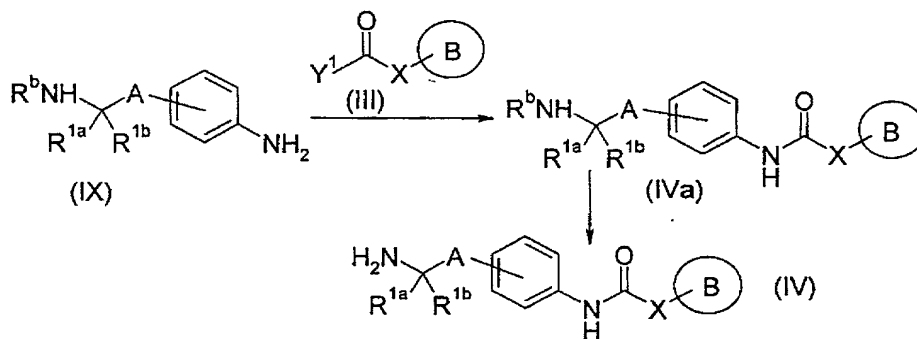
When R^b in the compound (VIIIa) is a hydrogen atom, the amino group is protected by customary manners using di-tert-butyl dicarbonate, etc., to prepare the compound (VIIIa).

Step (c):

A method for the reduction of nitro to amino may be conducted by customary manners such as metallic reduction using iron, zinc, etc. and catalytic reduction using a catalyst such as palladium-carbon, palladium hydroxide-carbon, Raney nickel, etc. R^a becomes a hydrogen atom depending upon the reduction conditions, but it may be protected again by customary manners.

(Manufacturing Method for Starting Compound (IV))

A)

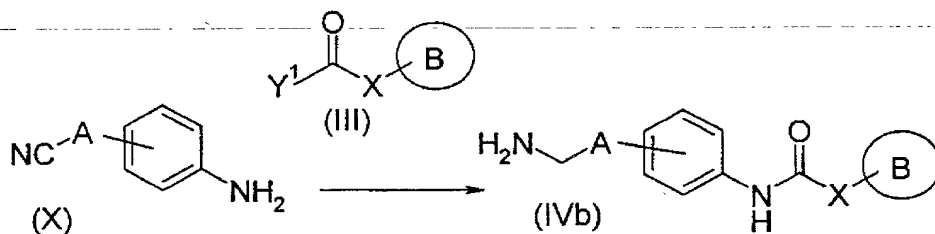


(In the formulae, R^{1a} , R^{1b} , R^b , A, B, X and Y^1 have the same meanings as defined already.)

This reaction is a reaction where the compound (IX) and the compound (III) are subjected to amidation reaction to give a compound (IVa) and, when R^b is a protective group for amino,

the protective group is removed to give a compound (IV). The amidation reaction can be conducted by the same manner as in the above-mentioned first manufacturing method, and the reaction conditions such as reaction temperature, solvent, etc. are the same as well.

B)



This reaction is a reaction where the compound (X) and the compound (III) are subjected to amidation reaction and then to reduction reaction to give a compound (IVb). The amidation reaction can be conducted by the same manner as in the above-mentioned first manufacturing method, and the reaction conditions such as reaction temperature, solvent, etc. are the same as well. In the reduction reaction, the above-mentioned catalytic reduction, or a method where reduction is conducted using sodium borohydride in the presence of cobalt chloride, may be applied.

With regard to other compounds such as the compound (III), the compound (V), the compound (VI), and the compound (VII), those which are available in the market or are appropriately synthesized by known methods (such as N-alkylation reaction, cyclization reaction, hydrolysis reaction, etc.) from the commercially available compounds may be used.

The compound (I) of the present invention which is manufactured as such is isolated and purified as a free compound, a salt thereof obtained by means of salt formation by customary manners, a hydrate, a solvate with various solvents such as ethanol, etc., or polymorphic crystals, etc. The isolation and purification may be conducted by applying common chemical operations such as extraction, concentration, evaporation, crystallization, filtration, recrystallization, various chromatographic methods, etc.

Various isomers may be isolated by customary manners utilizing the physico-chemical differences between the isomers. For example, the racemate can be converted to stereochemically pure isomers by common racemic resolution (such as a method where the racemate is changed to diastereomer salts with usual optically active acid (for example, tartaric acid), followed by optical resolution, and the like). Incidentally, a mixture of diastereomers may be separated by customary method such as fractional crystallization or chromatography, etc. In the case of an optically active

compound, it may be manufactured starting from an appropriate optically active material.

Industrial Applicability:

The phenethanol derivative of the present invention represented by the general formula (I) or the salt thereof has both an insulin secretion promoting action and an insulin sensitivity potentiating action and also has a selective β_3 -receptor stimulating action, so that it is useful as a therapeutic agent for diabetes mellitus.

As confirmed by a glucose tolerance test and a hypoglycemic test in insulin-resisting model animals as described later, the compound of the present invention has both a good insulin secretion promoting action and a good insulin sensitivity potentiating action, so that its usefulness in diabetes mellitus is expected. Although the β_3 -receptor stimulating action may have a possibility of participating in expression of the insulin secretion promoting action and the insulin sensitivity potentiating action, other mechanism might also possibly participate therein, and the details thereof have been still unknown yet. The β_3 -receptor stimulating action of the compound of the present invention is selective to β_3 -receptors in human being. It has been known that the stimulation of β_3 -receptor stimulates decomposition of fat (decomposition of the fat tissue triglyceride into

glycerol and free fatty acid), whereby a disappearance of fat mass is promoted. Therefore, the compound of the present invention has an anti-obesity action and an anti-hyperlipemia action (such as triglyceride lowering action, cholesterol lowering action and HDL cholesterol increasing action) and is useful as a preventive and therapeutic agent for obesity and hyperlipemia (such as hypertriglyceridemia, hypercholesterolemia and hypo-HDL-lipoproteinemia). Those diseases have been known as animus factors in diabetes mellitus, and amelioration of those diseases is useful for prevention and therapy of diabetes mellitus as well.

The compound of the present invention is also useful as a preventive and therapeutic agent for other diseases where the improvement of symptom can be achieved by reducing the symptoms of obesity and hyperlipemia such as ischemic coronary diseases such as arteriosclerosis, myocardial infarction, angina pectoris, etc. cerebral arteriosclerosis such as cerebral infarction, etc., or aneurysm, etc.

Further, the selective β_3 -receptor stimulating action of the compound of the present invention is useful for prevention and therapy of several diseases which have been reported to be improved by the stimulation of β_3 -receptor. Examples of those diseases are shown as follows.

It has been mentioned that the β_3 -receptor mediates the motility of non-sphincteral smooth muscle contraction, and

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because it is believed that the selective β_3 -receptor stimulating action assists the pharmacological control of intestinal motility without being accompanied by cardiovascular action, the compound of the present invention has a possibility of being useful in therapy of the diseases caused by abnormal intestinal motility such as various gastrointestinal diseases including irritable colon syndrome. It is also useful as the therapy for peptic ulcer, esophagitis, gastritis and duodenitis (including that induced by H. pylori), enterelcosis (such as inflammatory intestinal diseases, ulcerative colitis, clonal disease and proctitis).

It is further shown that the β_3 -receptor affects the inhibition of release of neuropeptide of some sensory fibers in lung. The sensory nerve plays an important role in neurogenic inflammation of respiratory tract including cough, and therefore, the specific β_3 -agonist of the present invention is useful in the therapy of neurogenic inflammation and in addition, has little action to cardiopulmonary system.

Moreover, the β_3 -adrenaline receptor is capable of resulting in a selective antidepressant action due to stimulation of the β_3 -receptor in brain, and accordingly, the compound of the present invention has a possibility of being useful as an antidepressant.

The action of the compound of the present invention has been ascertained to be selective to β_3 -receptors as a result

of experiments using cells expressing human type receptors, and the adverse action caused by other β_3 -receptor stimulation is low or none.

Effects of the compound of the present invention have been ascertained by the following tests.

1. Hypoglycemic test in kk mice (insulin-resisting model; obesity and hyperglycemia):

Male kk mice (blood sugar level: not lower than 200 mg/dl) were subjected to a measurement of blood sugar level under feeding and then randomly classified into groups. The drug to be tested was compulsorily administered orally or subcutaneously once daily for four days, and the blood sugar level after 15 to 18 hours from the final administration was compared with that before the administration (n = 6). The blood was collected from a tail vein of the mice using a glass capillary (previously treated with heparin), the protein was removed therefrom, and the amount of glucose in the supernatant liquid (mg/dl) was measured by colorimetric determination by means of a glucose oxidase method. Further, a dose at which the blood sugar level was lowered by 30% as compared with that before the administration with the drug to be tested was expressed as an ED₃₀ value.

As a result, the compound of the present invention significantly lowered the blood sugar level as compared with

that before the administration with the drug to be tested in both cases of oral and subcutaneous administrations. In particular, some of the compounds of the present invention exhibited a strong activity so that the ED₃₀ value in the oral administration was 3 mg/kg/day or less. On the other hand, in the above-referenced WO 95/29159, the compound of Example 90 had an ED₃₀ value of 30 mg/kg/day or more, and the compound of Example 92 had an ED₃₀ value of 30 mg/kg/day. From this fact, it has become clear that the compounds of the present invention have a superior potentiating action to insulin sensitivity as compared with those of the above-referenced WO 95/29159.

2. Glucose tolerance test in normal rats:

Male rats of SD strain of seven weeks age were fasted for a whole day and night, then randomly classified into groups and subjected to an oral glucose tolerance test (OGTT) (n = 4). The compound to be tested was administered orally or subcutaneously at 30 minutes before administration of glucose (2 g/kg by oral administration). The blood was collected from an abdominal aorta using a heparin-treated glass syringe from the rats which were anesthetized with pentobarbital (65 mg/kg), the protein was removed therefrom, and the amount of glucose in the supernatant liquid (mg/dl) was measured by colorimetric determination by means of a glucose oxidase method. The insulin value in blood was determined by measuring the amount

of insulin in plasma (ng/ml) by means of radioimmunoassay (RIA).

As a result, in a group where the compound of the present invention was administered orally or subcutaneously, a significant increase in the insulin value in blood was observed as compared with the group to which no drug was given. An increase in the sugar blood level after administration of glucose was significantly inhibited as well. From those results, it is apparent that the compound of the present invention has a good insulin secretion promoting action and a good hyperglycemia inhibiting action.

3. Stimulating test to human β_3 -, β_2 - and β_1 -receptors:

Human β_3 -stimulating action was investigated using an SK-N-MC cell system (cells in which human β_3 -receptor and human β_1 -receptor were permanently expressed were purchased) while human β_2 - and β_1 -stimulating actions were investigated using a CHO cell system (cells in which each of human β_2 - and β_1 -receptors was compulsorily expressed were purchased). Stimulating action of the compound (10^{-10} to 10^{-4} M) were investigated by incubating 10^5 cells/well of each of the cells on a 24-well plate and checking under a subconfluent state after two days using a producing activity of cyclic AMP (cAMP) as an index. Incidentally, the human β_3 -stimulating action was investigated in the presence of a β_1 -receptor blocker

(CGP20712A, 10^{-6} M). Amount of production of cAMP in each cell (pmol/ml) was measured by an RIA method using ^{125}I -cAMP. Intensity of action of each compound was compared by calculating the pD_2 value and the maximum activity (I.A. (%)) where the maximum reaction of 10^{-6} M isoproterenol was defined as 100%) from the resulting dose-reaction curve.

As a result, it has been ascertained that the compound of the present invention has a selective stimulating action to human β_3 -receptor.

A pharmaceutical composition containing one or more of the compound of the present invention or the salt thereof as an effective ingredient is prepared using common pharmaceutically acceptable vehicles. Administration of the pharmaceutical composition according to the present invention may be either by oral administration or by parenteral administration by, for example, injection, suppository, subcutaneous agent, inhaling agent or intracystic infusion.

The dose may be appropriately decided depending upon each particular case while taking into consideration symptom, age, sex, etc. of the patient but usually, is around 0.01 mg/kg to 100 mg/kg per day for adults in the case of oral administration, and that is administered at a time or by dividing into 2 to 4 times a day. When intravenous injection is conducted depending upon the symptom, the dose is usually around 0.001

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mg/kg to 10 mg/kg per day for adults, and that is administered at a time or by dividing into two or more times a day.

With regard to a vehicle for the preparation, nontoxic solid or liquid substances for pharmaceuticals may be used.

Examples of the solid composition for use by means of oral administration according to the present invention are tablets, pills, capsules, diluted powder and granules. In such a solid composition, one or more active substances are mixed with at least one inert excipient such as lactose, mannitol, glucose, hydroxypropyl cellulose, microcrystalline cellulose, starch, polyvinylpyrrolidone, agar, pectin, magnesium metasilicate aluminate and magnesium aluminate. The composition may also contain additives other than the inert excipient such as lubricants such as magnesium stearate; disintegrants such as calcium cellulose glycolate; stabilizers such as lactose; and auxiliary solubilizers such as glutamic acid or aspartic acid by customary manners. Tablets and pills may, if necessary, be coated with sugar coat such as sucrose, gelatin, hydroxypropyl cellulose, hydroxypropylmethyl cellulose phthalate, etc., or with film of gastric or enteric coating substances.

The liquid composition for oral administration includes pharmaceutically acceptable emulsions, solutions, suspensions, syrups and elixirs and contains commonly used inert excipients such as purified water or ethanol. In

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addition to the inert excipient, the composition may further contain auxiliary agents such as moisturizing or suspending agents, sweeteners, tasting agents, aromatic agents and antiseptic agents. The injection for parenteral administration includes aseptic aqueous or non-aqueous solutions, suspensions and emulsions. The non-aqueous solutions and suspensions include, for example, distilled water for injection and a physiological saline solution. Examples of the solvent for non-aqueous solution and suspension are propylene glycol; polyethylene glycol; plant oils such as cacao butter, olive oil and sesame oil; alcohols such as ethanol; gum arabic; and Polysolvate 80 (trade name). Such a composition may further contain auxiliary agents such as isotonizing agents; antiseptic agents; moisturizing agents; emulsifiers; dispersing agents; stabilizers such as lactose; and auxiliary solubilizers such as glutamic acid and aspartic acid). These may be sterilized, for example, by filtration passing through a bacteria-preserving filter or by compounding of or irradiation with a bactericide. These may also be used by manufacturing a sterile solid composition, followed by dissolving in sterile water or a sterile solvent for injection before use.

Best Mode for Carrying Out the Invention:

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The present invention is further illustrated by way of Examples as hereunder. Compounds of the present invention are not limited to those mentioned in the following Examples but cover all of the compounds represented by the above general formula (I), salts thereof, hydrates thereof, geometric and optical isomers thereof and polymorphic forms thereof. Incidentally, the case where the material which is used in the present invention is novel is illustrated by way of the following Referential Example.

Referential Example 1:

To a mixed solution of ethyl acetate and a 1N aqueous solution of sodium hydroxide was added 25.2 g of 4-nitrophenyl ethylamine hydrochloride, and the mixture was vigorously stirred. The organic layer was dried over anhydrous magnesium sulfate, and the solvent was evaporated. To the resulting residue were added 100 ml of 2-propanol and 15.0 g of (R)-styrene oxide successively, and the reaction mixture was heated to reflux for 12 hours. The solvent was evaporated in vacuo, and the residue was purified by silica gel column chromatography (eluent: chloroform/methanol = 100/1 → 10/1). The resulting residue was again subjected to silica gel column chromatography (eluent: hexane/ethyl acetate/triethylamine = 1/5/trace) to give 8.05 g of (R)-1-phenyl-2-[[2-(4-nitrophenyl)ethyl]amino]ethanol.

Referential Example 2:

A solution of 8.02 g of (R)-1-phenyl-2-[[2-(4-nitrophenyl)ethyl]amino]ethanol and 6.30 g of di-tert-butyl dicarbonate in 80 ml of tetrahydrofuran was stirred for 12 hours at room temperature. The residue obtained by evaporation of the solvent was purified by silica gel column chromatography (eluent: hexane/ethyl acetate = 3/1) to give 10.8 g of tert-butyl (R)-N-(2-hydroxy-2-phenylethyl)-N-[2-(4-nitrophenyl)ethyl]carbamate.

Referential Example 3:

To a solution of tert-butyl (R)-N-(2-hydroxy-2-phenylethyl)-N-[2-(4-nitrophenyl)ethyl]carbamate in 200 ml of ethanol was added 1.03 g of 10% palladium-carbon and the mixture was stirred for two hours at room temperature in a hydrogen atmosphere under atmospheric pressure. Insoluble matters were removed using Celite, and the filtrate was concentrated in vacuo to give 9.54 g of tert-butyl (R)-N-[2-(4-aminophenyl)-N-(2-hydroxy-2-phenylethyl)ethyl]-carbamate.

Referential Example 4:

To a solution of 448 mg of tert-butyl (R)-N-[2-(4-aminophenyl)-N-(2-hydroxy-2-phenylethyl)ethyl]carbamate

and 330 mg of triethylamine in 4 ml of chloroform was added 146 mg of 2-pyridinecarbonyl chloride. The reaction solution was stirred at room temperature for two hours, and the solvent was evaporated in vacuo. The residue was diluted with chloroform, and the organic layer was washed with a saturated aqueous solution of sodium hydrogen carbonate and dried over anhydrous magnesium sulfate. The residue obtained by evaporating the solvent in vacuo was purified by silica gel column chromatography (eluent: hexane/ethyl acetate = 1/3) to give 321 mg of tert-butyl (R)-N-(2-hydroxy-2-phenylethyl)-N-[2-[4-[(2-pyridinecarbonyl)amino]phenyl]ethyl]carbamate.

Referential Example 5:

To a solution of 377 mg of tert-butyl (R)-N-[2-(4-aminophenyl)-N-(2-hydroxy-2-phenylethyl)ethyl]carbamate in 10 ml of tetrahydrofuran were added 203 mg of 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide hydrochloride, 143 mg of 1-hydroxybenzotriazole and 202 mg of 8-quinolinecarboxylic acid successively. The reaction solution was stirred at room temperature for 18.5 hours, and the solvent was evaporated in vacuo. The residue was diluted with ethyl acetate, and the organic layer was washed with a saturated aqueous solution of sodium hydrogen carbonate and dried over anhydrous magnesium sulfate. The residue obtained by evaporation of the solvent was purified by silica gel column chromatography (eluent:

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hexane/ethyl acetate = 2/1) to give 302 mg of tert-butyl (R)-N-(2-hydroxy-2-phenylethyl)-N-[2-[4-[(8-quinolinecarbonyl)amino]phenyl]ethyl]carbamate.

Referential Example 6:

To a solution of 403 mg of tert-butyl (R)-N-(2-hydroxy-2-phenylethyl)-N-[2-[4-[(2-1H-imidazol-2-yl)acetyl]amino]phenyl]ethyl]carbamate in 10 ml of acetonitrile were added 120 mg of potassium carbonate and 164 mg of 2-fluorobenzyl bromide successively at room temperature. The reaction solution was stirred at 50°C for 12 hours. Insoluble matters were filtered off using Celite, and the solvent was evaporated. The resulting residue was purified by silica gel column chromatography to give 253 mg of tert-butyl (R)-N-[2-[4-[[2-[1-(2-fluorobenzyl)-1H-imidazol-2-yl]-acetyl]amino]phenyl]ethyl]-N-(2-hydroxy-2-phenylethyl)-carbamate.

Referential Example 7:

To a solution of 13.4 g of (R)-2-[N-benzyl-N-[2-(4-nitrophenyl)ethyl]amino]-1-phenylethanol in 150 ml of methanol were added 8.6 g of iron powder and 40 ml of a 2N aqueous hydrochloric acid solution. The reaction mixture was heated to reflux for two hours, a 1N aqueous solution of sodium hydroxide was added thereto, and the insoluble matters thus

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produced were filtered off using Celite. The filtrate was concentrated in vacuo to remove the methanol. The resulting aqueous phase was extracted with chloroform, the organic layer was dried over anhydrous magnesium sulfate, and the solvent was evaporated in vacuo. The resulting residue was purified by silica gel column chromatography (eluent: hexane/ethyl acetate = 1/1) to give 11.45 g of (R)-2-[N-[2-(4-amino-phenyl)ethyl]-N-benzylamino]-1-phenylethanol.

Referential Example 8:

To 502 mg of (R)-2-[N-[2-(4-aminophenyl)ethyl]-N-benzylamino]-1-phenylethanol were added 336 mg of ethyl 2-(3-methylpyridin-2-yl)acetate and 10 ml of xylene. The reaction mixture was refluxed for nine hours, and the solvent was evaporated in vacuo. The resulting residue was purified by silica gel column chromatography (eluent: hexane/ethyl acetate = 1/3) to give 222 mg of (R)-4'-[2-[N-benzyl-N-(2-hydroxy-2-phenylethyl)amino]ethyl]-2-(3-methylpyridin-2-yl)acetanilide.

Referential Example 9:

To a solution of 0.96 g of 2-fluoroacetophenone in 20 ml of tetrahydrofuran was added 2.65 g of benzyltrimethylammonium tribromide. The reaction mixture was stirred at room temperature for 30 minutes, insoluble matters were filtered

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off, and the solvent was concentrated in vacuo. The resulting residue was dissolved in 40 ml of 2-butanone, then 1.81 g of N-benzyl-4-nitrophenethylamine and 0.92 g of diisopropyl ethylamine were added, and the reaction mixture was heated to reflux for one hour. The solvent was evaporated in vacuo, ethyl acetate was added thereto, and the mixture was washed with water and a saturated saline solution successively. The organic layer was dried over anhydrous magnesium sulfate and evaporated in vacuo. The resulting residue was dissolved in 40 ml of methanol, 0.34 g of sodium borohydride was added thereto, and the reaction mixture was stirred at room temperature for one hour. The solvent was evaporated in vacuo, ethyl acetate was added, and the mixture was washed with water and a saturated saline solution successively. The organic layer was dried over anhydrous magnesium sulfate and evaporated in vacuo. The resulting residue was purified by silica gel column chromatography (eluent: chloroform) to give 1.95 g of 2-[N-benzyl-N-[2-(4-nitrophenyl)ethyl]amino]-1-(2-fluorophenyl)ethanol.

Referential Example 10:

A reaction mixture of 5.12 g of methyl 2-pyridylacetate, 5.14 g of 4-aminobenzyl cyanide and 50 ml of xylene was heated to reflux for 24 hours. An appropriate amount of the solvent was evaporated, diethyl ether was added to the residue, and

the resulting crystals were taken by filtration to give 5.65 g of 4'-cyanomethyl-2-(2-pyridyl)acetanilide.

Referential Example 11:

To a solution of 640 mg of 4'-cyanomethyl-2-(4,6-dimethyl-2-pyridyl)acetanilide in 15 ml of tetrahydrofuran was added 15 ml of an ethanolic suspension of a Raney nickel, and concentrated aqueous ammonia was added to adjust the pH of the mixture to about 10. The mixture was stirred at room temperature for one hour in a hydrogen atmosphere under atmospheric pressure. The reaction mixture was filtered using Celite, and the solvent was evaporated in vacuo to give 640 mg of 4'-(2-aminomethyl)-2-(4,6-dimethyl-2-pyridyl)acetanilide.

Referential Example 12:

To a solution of 630 mg of 4'-(2-aminomethyl)-2-(4,6-dimethyl-2-pyridyl)acetanilide in 20 ml of toluene was added 0.27 ml of benzaldehyde, and the mixture was heated to reflux for three hours using a Dean-Starke apparatus. The reaction mixture was filtered, and the solvent was evaporated in vacuo. A solution of the resulting residue in 30 ml of methanol was cooled at 0°C, 63 mg of sodium borohydride was added, and the mixture was stirred at 0°C for one hour. About one-half of the solvent of the reaction mixture was evaporated in vacuo, water

Example 1:

The compounds of Examples 2 to 33 were prepared by the same manner as in Example 1.

Example 2:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-3-pyridinecarboxanilide dihydrochloride

Example 3:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-8-quinolinecarboxanilide dihydrochloride

Example 4:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-(E)-3-(2-pyridyl)acrylic anilide dihydrochloride

Example 5:

(R)-2-(Benzothiazol-2-yl)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 6:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-(imidazo[2,1-b]thiazol-3-yl)acetanilide dihydrochloride

Example 7:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-(2-methylthiazol-4-yl)acetanilide hydrochloride

Example 8:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-(1H-imidazol-2-yl)acetanilide dihydrochloride

Example 9:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-(1H-tetrazol-5-yl)acetanilide hydrochloride

Example 10:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-(5-sulfanyl-1H-1,2,4-triazol-3-yl)acetanilide hydrochloride

Example 11:

(R)-2-(2-Aminothiazol-4-yl)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]-2-oxoacetanilide dihydrochloride

Example 12:

(R)-2-(5-Amino-1,2,4-thiadiazol-3-yl)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 13:

(R)-2-(5-Ethoxycarbonylamino-1,2,4-thiadiazol-3-yl)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide hydrochloride

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Example 14:

(R)-2-[(2-(3-Fluorophenylamino)thiazol-4-yl)-4']-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 15:

(R)-2-(2-Chloropyridin-6-yl)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide hydrochloride

Example 16:

(R)-2-(2-Benzyloxypyridin-6-yl)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide hydrochloride

Example 17:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-[1-(2-methyl-3-propenyl)-1H-imidazol-2-yl]acetanilide dihydrochloride

Example 18:

(R)-2-(1-Benzyl-1H-imidazol-4-yl)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 19:

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(R)-2-[1-(2-Chlorobenzyl)-1H-imidazol-4-yl]-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 20:

(R)-2-[1-(3-Chlorobenzyl)-1H-imidazol-4-yl]-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 21:

(R)-2-[1-(4-Chlorobenzyl)-1H-imidazol-4-yl]-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 22:

(R)-2-[1-(4-Fluorobenzyl)-1H-imidazol-2-yl]-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 23:

(R)-2-[1-(4-Chlorobenzyl)-1H-imidazol-2-yl]-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 24:

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(R)-2-[1-(4-Bromobenzyl)-1H-imidazol-2-yl]-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 25:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-[1-(4-iodobenzyl)-1H-imidazol-2-yl]acetanilide dihydrochloride

Example 26:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-[1-(4-trifluoromethylbenzyl)-1H-imidazol-2-yl]acetanilide dihydrochloride

Example 27:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-[1-(2-naphthyl)-1H-imidazol-2-yl]acetanilide dihydrochloride

Example 28:

(R)-2-[1-(4-Fluorobenzyl)-5-methyl-1H-imidazol-2-yl]-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 29:

(R)-2-[1-(4-Fluorobenzyl)-4-methyl-1H-imidazol-2-yl]-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

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Example 30:

(R)-2-[1-(4-Fluorobenzyl)-1H-tetrazol-5-yl]-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide hydrochloride

Example 31:

(R)-2-[2-(3,4-Dichlorobenzyl)-1H-tetrazol-5-yl]-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide hydrochloride

Example 32:

(R)-2-[2-(4-Fluorobenzyl)-1H-tetrazol-5-yl]-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide hydrochloride

Example 33:

(R)-2-[1-(3,4-Dichlorobenzyl)-1H-tetrazol-5-yl]-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide hydrochloride

Example 34:

To a solution of 175 mg of tert-butyl (R)-N-[2-[4-[2-(1H-1,2,4-triazol-3-yl)acetylamino]phenyl]ethyl]-N-(2-hydroxy-2-phenylethyl)carbamate in 5 ml of methanol was added

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4 ml of a solution of 4N hydrogen chloride in ethyl acetate. The mixture was stirred at room temperature for three hours, the solvent was filtered off, and the resulting powder was washed with ethanol. The resulting powder was dried to give 125 mg of (R)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]-2-(1H-1,2,4-triazol-3-yl)acetanilide dihydrochloride.

The compounds of Examples 35 to 40 were prepared by the same manner as in Example 34.

Example 35:

(R)-2-(5-Benzylsulfanyl-1H-1,2,4-triazol-3-yl)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 36:

(R)-2-(2-Acetamidothiazol-4-yl)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide hydrochloride

Example 37:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-(2-methanesulfonamidothiazol-4-yl)acetanilide hydrochloride

Example 38:

(R)-2-(2-Guanidinothiazol-4-yl)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

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Example 39:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-(2-phenylaminothiazol-4-yl)acetanilide hydrochloride

Example 40:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-[1-(4-nitrobenzyl)-1H-imidazol-2-yl]acetanilide hydrochloride

Example 41:

To 690 mg of tert-butyl (R)-N-[2-[4-[2-(2-aminothiazol-4-yl)acetamino]phenyl]ethyl]-N-[(2-hydroxy-2-phenyl)ethyl]carbamate were added 30 ml of methanol and 15 ml of a solution of 4N hydrogen chloride in ethyl acetate, and the mixture was stirred at room temperature for two hours. The solvent was evaporated in vacuo, and the residue was purified by a reverses phase column chromatography (eluent: water/methanol = 2/1) to give 310 mg of (R)-2-(2-aminothiazol-4-yl)-4'-[2-(2-hydroxy-2-phenylethyl)amino]-ethyl]acetanilide dihydrochloride.

The compounds of Examples 42 to 57 were prepared by the same manner as in Example 41.

Example 42:

Example 43:

Example 44:

Example 45:

Example 46:

Example 47:

Example 48:

(R)-2-(1-Benzyl-1H-1,2,4-triazol-3-yl)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide hydrochloride

Example 49:

(R)-2-(3-Benzyl-2-thioxothiazol-4-yl)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide hydrochloride

Example 50:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-(5,6,7,8-tetrahydroquinolin-8-yl)carboxanilide dihydrochloride

Example 51:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-(1-phenyl-1H-imidazol-2-yl)acetanilide dihydrochloride

Example 52:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-[(1-(4-isopropylbenzyl)-1H-imidazol-2-yl)acetanilide dihydrochloride

Example 53:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-[(1-(4-phenylbenzyl)-1H-imidazol-2-yl)acetanilide dihydrochloride

Example 54:

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(R)-2-[1-(2-Chlorobenzyl)-1H-imidazol-2-yl]-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 55:

(R)-2-[1-(3-Chlorobenzyl)-1H-imidazol-2-yl]-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 56:

(R)-2-[1-(3,4-Dichlorobenzyl)-1H-imidazol-2-yl]-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 57:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-[(1-(2-pyridyl)methyl-1H-imidazol-2-yl)acetanilide dihydrochloride

The compound of Example 58 was prepared by the same manner as in Example 1.

Example 58:

(R)-2-(2-aminopyridin-6-yl)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

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Example 59:

To a solution of tert-butyl (R)-N-[2-[4-[[2-(2-aminothiazol-4-yl)-2-oxoacetyl]amino]phenyl]ethyl]-N-(2-hydroxy-2-phenylethyl)carbamate in 30 ml of methanol was added 130 mg of sodium borohydride at room temperature. The reaction mixture was stirred at room temperature for three hours, and the solvent was evaporated in vacuo. The residue was dissolved in 5 ml of methanol, and to this reaction solution was added 10 ml of a solution of 4N hydrogen chloride-ethyl acetate. The reaction solution was stirred at room temperature for eight hours and the solvent was evaporated in vacuo. The residue was purified by silica gel column chromatography (eluent: chloroform/methanol = 5/1). The resulting residue was purified by reversed phase column chromatography (eluent: water/methanol = 2/1) to give 77 mg of (R)-2-(2-aminothiazol-4-yl)-2-hydroxy-4'-[2-(2-hydroxy-2-phenylethyl)-amino]acetanilide hydrochloride.

Example 60:

To 349 mg of tert-butyl (R)-N-[2-[4-[[2-(2-benzoyloxy)pyridin-6-yl]acetyl]amino]phenyl]ethyl]-N-(2-hydroxy-2-phenylethyl)carbamate were added 478 mg of pentamethylbenzene and 5 ml of trifluoroacetic acid successively. The reaction solution was stirred at room temperature for four hours, and the solvent was evaporated in vacuo. To the residue were added

water and potassium carbonate to make the solution basic, and the aqueous phase was extracted with a mixed solvent of chloroform and tetrahydrofuran. The organic layer was dried over anhydrous magnesium sulfate, and the solvent was evaporated in vacuo. The residue was purified by silica gel column chromatography (eluent: chloroform/methanol = 10/1 → 5/1). To an ethanolic solution of the resulting residue was added 100 µl of a 4N hydrogen chloride-ethyl acetate solution, and then the solvent was evaporated in vacuo. The resulting crude crystals were recrystallized from ethanol-ethyl acetate to give 65 mg of (R)-2-(2-benzyloxypyridin-6-yl)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide hydrochloride.

The compounds of Examples 61 to 76, 83 and 85 were prepared by the same manner as in Example 1; and the compounds of Examples 77 to 82 were prepared by the same manner as in Example 41.

Example 61:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-(2-methylpropyl-1H-imidazol-2-yl)acetanilide dihydrochloride

Example 62:

(R)-2-[1-(2-Fluorobenzyl)-1H-imidazol-2-yl]-4'-[2-(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 63:

(R)-2-[1-(3-Fluorobenzyl)-1H-imidazol-2-yl]-4'-[2-(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 64:

(R)-2-[1-(2,4-Difluorobenzyl)-1H-imidazol-2-yl]-4'-[2-(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 65:

(R)-2-[1-(2,6-Difluorobenzyl)-1H-imidazol-2-yl]-4'-[2-(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 66:

(R)-2-[1-(3,5-Difluorobenzyl)-1H-imidazol-2-yl]-4'-[2-(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 67:

(R)-2-[1-(2,5-Difluorobenzyl)-1H-imidazol-2-yl]-4'-[2-(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 68:

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(R)-2-[1-(3,4-Difluorobenzyl)-1H-imidazol-2-yl]-4'-[2-(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 69:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-[1-(2,3,6-trifluorobenzyl)-1H-imidazol-2-yl]acetanilide dihydrochloride

Example 70:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-[1-(2,4,5-trifluorobenzyl)-1H-imidazol-2-yl]acetanilide dihydrochloride

Example 71:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-[1-(3,4,5-trifluorobenzyl)-1H-imidazol-2-yl]acetanilide dihydrochloride

Example 72:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-[1-(2,3,4,5,6-pentafluorobenzyl)-1H-imidazol-2-yl]acetanilide dihydrochloride

Example 73:

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(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-[1-(3-iodobenzyl)-1H-imidazol-2-yl]acetanilide dihydrochloride

Example 74:

(R)-2-[1-(2,6-Dichlorobenzyl)-1H-imidazol-2-yl]-4'-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide hydrochloride

Example 75:

(R)-2-[1-(4-Cyanobenzyl)-1H-imidazol-2-yl]-4'-[2-(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 76:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-[1-(quinolin-2-yl)-1H-imidazol-2-yl]acetanilide trihydrochloride

Example 77:

(R)-2-[1-(2-Chloro-6-fluorobenzyl)-1H-imidazol-2-yl]-4'-[2-(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide

Example 78:

(R)-2-[1-(2-Chloro-4-fluorobenzyl)-1H-imidazol-2-yl]-4'-[2-(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide

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Example 79:

(R)-2-[1-(2,5-Dichlorobenzyl)-1H-imidazol-2-yl]-4'-[2-(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 80:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-[1-(2,3,4-trifluorobenzyl)-1H-imidazol-2-yl]acetanilide dihydrochloride

Example 81:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-[1-(4-methoxycarbonylbenzyl)-1H-imidazol-2-yl]acetanilide dihydrochloride

Example 82:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-[1-[(piperidine-1-carbonyl)benzyl]-1H-imidazol-2-yl]acetanilide dihydrochloride

Example 83:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-(1-pyrazolyl)acetanilide hydrochloride

Example 84:

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(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-(1,2,4-triazol-1-yl)acetanilide dihydrochloride

Example 85:

(R)-2-(2-Aminobenzimidazol-1-yl)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 86:

To a solution of 20.1 g of 4'-[2-[N-benzyl-N-(2-hydroxy-2-phenylethyl)amino]ethyl]-2-(2-pyridyl)acetanilide in 400 ml of methanol was added 5.96 g of 10% palladium-carbon. The reaction solution was stirred for six hours in a hydrogen atmosphere under atmospheric pressure. Insoluble matters were filtered off using Celite and the filtrate was concentrated in vacuo. To a methanolic solution of the resulting residue was added 10.8 ml of a 4N hydrogen chloride-ethyl acetate solution, and the solvent was evaporated in vacuo. The resulting crude crystals were recrystallized from methanol-ethanol to give (R)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]-2-(2-pyridyl)acetanilide hydrochloride.

The compounds of 87 to 90 were prepared by the same manner as in Example 86.

Example 87:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-(3-pyridyl)acetanilide hydrochloride

Example 88:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-(4-pyridyl)acetanilide hydrochloride

Example 89:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-3-(2-pyridyl)propionanilide hydrochloride

Example 90:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-[(1-phenylethyl)-1H-imidazol-2-yl]acetanilide dihydrochloride

Example 91:

(R)-2-(1H-Benzimidazol-2-yl)-4'-[4-[2-[N-benzyl-N-(2-hydroxy-2-phenylethyl)amino]ethyl]phenyl]acetanilide (240 mg) was dissolved in 30 ml of ethanol, then 170 mg of 10% palladium-carbon was added thereto and the mixture was stirred for nine hours in a hydrogen atmosphere under atmospheric pressure. The catalyst was filtered off, the solvent was evaporated in vacuo, and the residue was washed with ethanol-ethyl acetate to give 200 mg of (R)-2-(1H-

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benzimidazol-2-yl)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]-ethyl]acetanilide.

The compounds of Examples 92 and 93 were prepared by the same manner as in Example 86.

Example 92:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-(3-methylpyridin-2-yl)acetanilide hydrochloride

Example 93:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-(2-pyrazinyl)acetanilide hydrochloride

Example 94:

(R)-4'-[4-[2-[N-Benzyl-N-(2-hydroxy-2-phenylethyl)-amino]ethyl]phenyl]-2-(1-benzyl-1H-imidazol-2-yl)acetanilide (350 mg) was dissolved in 20 ml of ethanol, then 130 mg of 10% palladium-carbon was added thereto, and the mixture was stirred for 17.5 hours in a hydrogen atmosphere under atmospheric pressure. The catalyst was filtered off, the solvent was evaporated in vacuo, and the residue was purified by silica gel column chromatography (eluent: chloroform/methanol/concentrated aqueous ammonia = 200/10/1). The resulting oily substance was dissolved in methanol, and 280 μ l of a 4N hydrogen chloride-ethyl acetate solution was added

thereto. The mixture was filtered after adding active carbon was added thereto, and the solvent was evaporated in vacuo to give 200 mg of (R)-2-(1-benzyl-1H-imidazol-2-yl)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride.

The compounds of Examples 95 and 97 were prepared by the same manner as in Example 91; the compounds of Examples 98 and 100 were prepared by the same manner as in Example 94; and the compounds of Examples 99 and 101 to 103 were prepared by the same manner as in Example 86.

Example 95:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-(4-methyl-2-pyridyl)acetanilide

Example 96:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-(5-methyl-2-pyridyl)acetanilide

Example 97:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-(6-methyl-2-pyridyl)acetanilide

Example 98:

4'-[(R)-2-[(R)-2-Hydroxy-2-phenylethyl]amino]propyl]-2-(2-pyridyl)acetanilide hydrochloride

Example 99:

4'-[(S)-2-[(R)-2-Hydroxy-2-phenylethyl]amino]propyl]-2-(2-pyridyl)acetanilide hydrochloride

Example 100:

2-(1-Benzyl-1H-imidazol-2-yl)-4'-[(S)-2-[(R)-2-hydroxy-2-phenylethyl]amino]propyl]acetanilide hydrochloride

Example 101:

4'-[2-[[2-Hydroxy-2-(2-fluorophenyl)ethyl]amino]ethyl]-2-(2-pyridyl)acetanilide hydrochloride

Example 102:

4'-[2-[[2-Hydroxy-2-(3-fluorophenyl)ethyl]amino]ethyl]-2-(2-pyridyl)acetanilide hydrochloride

Example 103:

4'-[2-[[2-Hydroxy-2-(4-fluorophenyl)ethyl]amino]ethyl]-2-(2-pyridyl)acetanilide hydrochloride

Example 104:

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To a solution of 805 mg of 4'-cyanomethyl-2-(2-pyrimidinyl)acetanilide in 30 ml of tetrahydrofuran were added 30 ml of an ethanolic solution of a Raney nickel and 3 ml of concentrated aqueous ammonia. The reaction solution was stirred for four hours in a hydrogen atmosphere under atmospheric pressure, then insoluble matters were filtered off using Celite, and the solvent was evaporated. To the resulting residue were added 10 ml of 2-propanol, 300 mg of (R)-styrene oxide and 2 ml of methanol successively. The reaction mixture was heated to reflux for ten hours, and the solvent was evaporated. The residue was purified by silica gel column chromatography (eluent: chloroform/methanol = 10/1). To a methanolic solution of the resulting residue was added 150 μ l of 4N hydrogen chloride-ethyl acetate solution, and the solvent was evaporated in vacuo. The resulting residue was crystallized from methanol-ethanol-ethyl acetate and then recrystallized from ethanol-diethyl ether to give 160 mg of (R)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]-2-(2-pyrimidinyl)acetanilide hydrochloride.

The compounds of Examples 105 to 108 were prepared by the same manner as in Example 104; and the compound of Example 109 was prepared by the same manner as in Example 91.

Example 105:

(R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-(2-quinolyl)acetanilide hydrochloride

Example 106:

(R)-4'-[2-[[2-Hydroxy-2-(3-chlorophenyl)ethyl]amino]-ethyl]-2-(2-pyridyl)acetanilide hydrochloride

Example 107:

4'-[2-[[2-Hydroxy-2-(3-pyridyl)ethyl]amino]ethyl]-2-(2-pyridyl)acetanilide hydrochloride

Example 108:

(R)-2-[1-(4-Chlorobenzyl)-1H-benzimidazol-2-yl]-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide dihydrochloride

Example 109:

(R)-2-(4,6-Dimethyl-2-pyridyl)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide

Example 110:

To 4'-(3-aminopropyl)-2-(2-pyridyl)acetanilide were added 10 ml of 2-propanol and 600 mg of (R)-styrene oxide successively. The reaction mixture was heated to reflux for four hours, and the solvent was evaporated. The residue was

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purified by silica gel column chromatography (eluent: chloroform/methanol = 30/1 \rightarrow 10/1). To a methanolic solution of the resulting residue was added 100 μ l of a 4N hydrogen chloride-ethyl acetate solution, and the solvent was evaporated in vacuo. The resulting crude crystals were recrystallized from ethanol-diethyl ether to give 71 mg of (R)-4'-[3-[(2-hydroxy-2-phenylethyl)amino]propyl]-2-(2-pyridyl)acetanilide hydrochloride.

Example 111:

To a solution of 3.62 g of tert-butyl N-[2-[4-[[2-(2-pyridyl)acetyl]amino]phenoxy]ethyl]carbamate in 30 ml of methanol was added 50 ml of a 4N hydrochloride-ethyl acetate solution. After the reaction solution was stirred at room temperature for eight hours, the solvent was evaporated in vacuo. To the residue were added an aqueous solution of sodium hydrogen carbonate and potassium carbonate to adjust to pH about 12. The resulting aqueous phase was extracted with a mixed solvent of chloroform and tetrahydrofuran. The organic layer was dried over anhydrous magnesium sulfate and concentrated, the resulting residue was dissolved in 40 ml of methanol, and 1.02 g of (R)-styrene oxide was added thereto. After the reaction solution was heated to reflux for 26 hours, the solvent was evaporated in vacuo. The resulting residue was purified by silica gel column chromatography (eluent:

chloroform/methanol = 30/1 \rightarrow 10/1) and dissolved in methanol, 0.59 ml of a 4N hydrogen chloride-ethyl acetate solution was added, and the solvent was evaporated in vacuo. The resulting crude crystals were recrystallized from methanol-ethanol to give 320 mg of (R)-4'-[2-[(2-hydroxy-2-phenylethyl)-amino]ethoxy]-2-(2-pyridyl)acetanilide hydrochloride

Example 112:

To a solution of 490 mg of tert-butyl N-[1,1-dimethyl-2-[4-[[2-(2-pyridyl)acetyl]amino]phenyl]ethyl]-carbamate in 10 ml of methanol was added 30 ml of a 4N hydrochloride-ethyl acetate solution. After the reaction solution was stirred at room temperature for eight hours, the solvent was evaporated in vacuo. To the residue were added an aqueous solution of sodium hydrogen carbonate and potassium carbonate to adjust to pH about 12. The resulting aqueous phase was extracted with a mixed solvent of chloroform and tetrahydrofuran. The organic layer was dried over anhydrous magnesium sulfate and concentrated, the resulting residue was dissolved in 2 ml of 2-propanol and 2 ml of methanol, and 120 mg of (R)-styrene oxide was added thereto. After the reaction solution was heated to reflux for 24 hours, the solvent was evaporated in vacuo. The resulting residue was purified by silica gel column chromatography (eluent: chloroform/methanol = 30/1 \rightarrow 5/1) and dissolved in methanol, 0.1 ml of a 4N hydrogen

chloride-ethyl acetate solution was added, and the solvent was evaporated in vacuo. The resulting residue was purified by silica gel column chromatography (eluent: chloroform/methanol = 5/1) and a reversed phase column chromatography (eluent: water/methanol = 2/1 → 1/1) to give 35 mg of (R)-4'-[2,2-dimethyl-2-[(2-hydroxy-2-phenylethyl)amino]ethyl]-2-(2-pyridyl)acetanilide hydrochloride.

The compound of Example 113 was prepared by the same manner as in Example 1.

Example 113:

(R)-1-[4-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]phenyl]-3-(2-pyridyl)urea dihydrochloride

As hereunder, physical and chemical properties of the compounds of the Referential Examples are given in Tables 1 and those of the compounds of the Examples are given in Tables 2.

The symbols in the tables have the following meanings.

Rex.: Referential Example No.

Ex.: Example No.

DATA: Physico-chemical properties

NMR: Nucleomagnetic resonance spectrum (TMS internal standard; DMSO-d was used as a solvent unless otherwise specified)

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mp: melting point

dec: decomposition

MS (m/z): mass spectrographic data (m/z)

Structure: structural formula

Table 1

Rex	DATA
1	NMR (CDCl ₃) δ : 2.75(1H,dd,J=12.4, 8.8Hz), 2.85-3.04(5H,m), 4.70(1H,dd,J=8.8, 3.7Hz), 7.24-7.40(7H,m), 8.10-8.20(2H,m)
2	NMR (CDCl ₃) δ : 1.44(9H,s), 2.75-3.10(2H,m), 3.20-3.70(4H,m), 4.93(1H,br), 7.25-7.40(7H,m), 8.14(2H,d,J=8.4 Hz)
3	NMR (CDCl ₃) δ : 1.47(9H,s), 2.55-2.80(2H,m), 3.20-3.40(2H,m), 3.45-3.65(2H,m), 4.87(1H,m), 6.57-6.65(2H,m), 6.83-7.04(2H,m), 7.25-7.40(5H,m)
4	NMR (CDCl ₃) δ : 1.47(9H,s), 2.62-2.93(2H,m), 3.14-3.58(4H,m), 4.35(1H,brs), 4.90(1H,br), 7.06-7.40(7H,m), 7.45-7.50(1H,m), 7.67-7.72(2H,m), 7.90(1H,dt,J=2.0, 8.0Hz), 8.25-8.31(1H,m), 8.58-8.63(1H,m), 9.98(1H,brs)
5	NMR (CDCl ₃) δ : 1.49(9H,s), 2.64-2.90(2H,m), 3.16-3.60(4H,m), 4.38(1H,brs), 4.91(1H,br), 7.10-7.42(7H,m), 7.55(1H,dd,J=8.0, 4.4Hz), 7.74(1H,t,J=8.0Hz), 7.77-7.84(2H,m), 8.01(1H,d,J=8.0, 1.2Hz), 8.34(1H,d,J=8.4, 1.6Hz), 8.96(1H,d,J=7.6, 1.6Hz), 9.02(1H,d,J=4.4, 2.0Hz), 13.61(1H,brs)
6	NMR (CDCl ₃) δ : 1.47(9H,s), 2.60-2.80(2H,m), 3.15-3.55(4H,m), 3.78(2H,s), 4.36(1H,brs), 4.82-4.94(1H,m), 5.18(2H,s), 6.92-6.99(2H,m), 7.00-7.13(5H,m), 7.25-7.38(6H,m), 7.42-7.48(2H,m), 10.34(1H,brs)
7	NMR (CDCl ₃) δ : 2.56-2.94(6H,m), 3.40-3.65(2H,m), 3.80(1H,brs), 3.95(1H,d,13.6Hz), 4.62(1H,dd,J=10.0, 3.2Hz), 6.57-6.66(2H,m), 6.87-6.98(2H,m), 7.20-7.37(10H,m)
8	NMR (CDCl ₃) δ : 2.40(3H,s), 2.54-3.00(6H,m), 3.57(1H,d,J=13.6Hz), 3.88(2H,s), 3.95(1H,d,J=13.6Hz), 4.62(1H,dd,J=10.4, 3.6Hz), 7.00-7.75(16H,m), 8.44(1H,d,J=4.4Hz), 9.66(1H,brs)
9	NMR (CDCl ₃) δ : 2.58-2.65(1H,m), 2.75-3.00(5H,m), 3.59(1H,d,J=13.2Hz), 3.95(1H,d,J=13.2Hz), 5.01(1H,dd,J=10.0, 3.2Hz), 6.97-7.03(1H,m), 7.12-7.35(9H,m), 7.48-7.56(1H,m), 8.04-8.13(2H,m)
10	NMR (CDCl ₃) δ : 3.70(2H,s), 3.88(2H,s), 7.23-7.32(4H,m), 7.54-7.62(2H,m), 7.71(1H,dt,J=7.6, 1.6Hz), 8.63(1H,d), 10.04(1H,brs)
11	NMR (CDCl ₃) δ : 2.26(3H,s), 2.39(3H,s), 2.57(2H,t,J=7.2Hz), 2.72(2H,t,J=7.2Hz), 3.72(2H,s), 6.95(1H,s), 7.01(1H,s), 7.11(2H,d,J=8.8Hz), 7.51(2H,d,J=8.8Hz), 10.17(1H,s)
12	NMR δ : 2.32(3H,s), 2.41(3H,s), 2.90-3.19(6H,m), 3.75(2H,s), 4.01(2H,s), 4.89(1H,dt,J=7.6, 3.2Hz), 6.99-7.71(16H,m), 10.26(1H,s)

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Table 2

Ex	D A T A
1	mp: 223-225°C. NMR δ : 2.95-3.28(6H,m), 4.98-5.07(1H,m), 7.23-7.44(6H,m), 7.65-7.75(1H,m), 7.88(2H,d,J=8.4Hz), 8.05-8.22(2H,m), 8.75(1H,d,J=4.4Hz), 8.97(1H,brs), 9.43(1H,brs), 10.65(1H,brs)
2	mp: 263-265°C. NMR δ : 2.92-3.10(3H,m), 3.13-3.27(3H,m), 5.00(1H,dd,J=10.8, 2.8Hz), 7.24-7.44(8H,m), 7.74-7.81(3H,m), 8.57(1H,d,J=8.0Hz), 8.81-8.96(2H,m), 9.20-9.30(2H,m), 10.71(1H,brs)
3	mp: 145-147°C. NMR δ : 2.94-3.10(3H,m), 3.14-3.30(3H,m), 4.97-5.05(1H,m), 7.27-7.46(7H,m), 7.77-7.90(4H,m), 8.30(1H,dd,J=8.4, 1.6Hz), 8.60-8.71(2H,m), 8.89(1H,brs), 9.10-9.30(2H,m), 13.12(1H,brs)
4	mp: 246-248°C (dec). NMR δ : 2.92-3.09(3H,m), 3.11-3.26(3H,m), 5.01(1H,dd,J=10.4, 2.8Hz), 7.24(2H,d,J=8.4Hz), 7.29-7.47(6H,m), 7.56-7.75(4H,m), 7.85(1H,d,J=8.0Hz), 8.11(1H,t,J=7.6Hz), 8.73(1H,d,J=4.4Hz), 8.92(1H,brs), 9.32(1H,brs), 10.69(1H,brs)
5	mp: 228-233°C (dec). NMR δ : 2.88-3.09(3H,m), 3.10-3.24(3H,m), 4.30(2H,s), 4.93-5.01(1H,m), 6.19(1H,d,J=3.6Hz), 7.18-7.27(2H,m), 7.28-7.53(7H,m), 7.57-7.62(2H,m), 7.97(1H,d,J=7.6Hz), 8.08(1H,d,J=8.0Hz), 8.83(1H,brs), 9.11(1H,brs), 10.57(1H,brs)
6	mp: 161-162°C. NMR δ : 2.86-3.24(6H,m), 4.24(2H,s), 4.97(1H,dd,J=9.6, 2.8Hz), 7.16-7.23(2H,m), 7.27-7.44(5H,m), 7.55(1H,s), 7.61(2H,d,J=8.4Hz), 7.85(1H,s), 8.27(1H,d,J=2.4Hz), 8.97(1H,brs), 9.47(1H,brs), 10.94(1H,brs)
7	NMR δ : 2.70(3H,s), 2.86-3.27(6H,m), 3.85(2H,s), 5.00-5.05(1H,m), 7.18-7.60(10H,m), 10.43(1H,s)
8	mp: 203-207°C. NMR δ : 2.92-3.08(3H,m), 3.10-3.22(3H,m), 4.28(2H,s), 5.01(1H,d,J=7.8Hz), 6.21(1H,brs), 7.22(2H,d,J=8.3Hz), 7.25-7.63(4H,m), 8.93(1H,brs), 9.38(1H,brs), 10.86(1H,s)
9	mp: 259-261°C. NMR δ : 2.90-3.10(3H,m), 3.10-3.25(3H,m), 4.15(2H,s), 4.97(1H,d,J=10.8Hz), 6.20(1H,d,J=3.9Hz), 7.21(2H,d,J=8.8Hz), 7.30-7.42(5H,m), 7.57(2H,d,J=8.8Hz), 8.85(1H,brs), 9.14(1H,brs), 10.58(1H,s)
10	mp: 210-213°C. NMR δ : 2.86-3.08(3H,m), 3.12-3.22(3H,m), 3.73(2H,s), 4.91-4.98(1H,m), 6.19(1H,d,J=3.9Hz), 7.21(2H,d,J=8.3Hz), 7.29-7.42(5H,m), 7.54(2H,d,J=8.3Hz), 8.78(1H,brs), 8.99(1H,brs), 10.35(1H,s), 13.21(1H,brs), 13.34(1H,brs)
11	mp: 205-210°C (dec). NMR δ : 2.90-3.25(6H,m), 4.95-5.04(1H,m), 7.23-7.44(7H,m), 7.67-7.75(2H,m), 8.15(1H,s), 8.88(1H,brs), 9.25(1H,brs), 10.83(1H,brs)
12	mp: 244-246°C. NMR δ : 2.90-3.08(3H,m), 3.10-3.20(3H,m), 3.67(2H,s), 5.00(1H,dd,J=2.4, 10.02Hz), 7.19(2H,d,J=8.3Hz), 7.28-7.42(5H,m), 7.57(2H,d,J=8.3Hz), 8.90(1H,s), 9.31(1H,s), 10.31(1H,s)
13	mp: 205-208°C. NMR δ : 1.27(3H,t,J=7.1Hz), 2.88-3.08(3H,m), 3.12-3.22(3H,m), 3.86(2H,s), 4.27(2H,q,J=7.1Hz), 4.96(1H,d,J=8.3Hz), 6.20(1H,s), 7.19(2H,d,J=8.3Hz), 7.30-7.42(5H,m), 7.57(2H,d,J=8.3Hz), 8.81(1H,s), 9.10(1H,s), 10.33(1H,s), 12.53(1H,s)
14	mp: 169-173°C. NMR δ : 2.88-3.22(6H,m), 3.66(2H,s), 4.98(1H,dd,J=2.9, 13.1Hz), 6.72(1H,s), 7.19(2H,d,J=8.3Hz), 7.23-7.42(8H,m), 7.59(2H,d,J=8.3Hz), 7.72-7.78(1H,m), 8.85(1H,s), 9.18(1H,brs), 10.24(1H,brs), 10.55(1H,s)
15	mp: 248-251°C. NMR δ : 2.90-3.08(3H,m), 3.09-3.21(3H,m), 3.88(2H,s), 5.02(1H,dd,J=10.0, 2.4Hz), 6.20(1H,brs), 7.16-7.22(2H,m), 7.28-7.46(7H,m), 7.57-7.63(2H,m), 7.84(1H,t,J=7.2Hz), 8.95(1H,brs), 9.40(1H,brs), 10.48(1H,brs)
16	mp: 237-238°C. NMR δ : 2.87-3.24(6H,m), 3.77(2H,s), 4.93-5.03(1H,m), 5.32(2H,s), 6.20(1H,d,J=4.0Hz), 6.73(1H,d,J=8.0Hz), 6.99(1H,d,J=7.2Hz), 7.16-7.22(2H,m), 7.25-7.46(10H,m), 7.57-7.63(2H,s), 7.67(1H,dd,J=8.4, 7.2Hz), 8.87(1H,brs), 9.24(1H,brs), 10.30(1H,brs)
17	mp: 190-193°C. NMR δ : 1.68(3H,m), 2.90-3.10(3H,m), 3.10-3.20(3H,m), 4.32(2H,s), 4.67(1H,s), 4.83(2H,s), 4.94(1H,s), 4.99(1H,d,J=8.3Hz), 6.21(1H,brs), 7.21(2H,d,J=8.7Hz), 7.24-7.42(5H,m), 7.56(2H,d,J=8.8Hz), 7.66(2H,d,J=1.9Hz), 7.71(1H,d,J=1.9Hz), 8.89(1H,brs), 9.30(1H,brs), 10.92(1H,s)
18	mp: 139-141°C. NMR δ : 3.01(3H,brs), 3.15(3H,brs), 3.92(2H,s), 5.05(1H,d,J=10.3Hz), 5.44(2H,s), 6.19(1H,brs), 7.19(2H,d,J=8.3Hz), 7.31-7.47(10H,m), 7.60(2H,d,J=8.3Hz), 7.66(1H,s), 9.05(1H,brs), 9.35(1H,s), 9.60(1H,brs), 10.76(1H,s)
19	mp: 140-143°C. NMR δ : 2.99-3.09(3H,m), 3.16(3H,brs), 3.95(2H,s), 5.06(1H,d,J=10.4Hz), 5.57(2H,s), 6.19(1H,brs), 7.19(2H,d,J=8.6Hz), 7.29-7.35(1H,m), 7.37-7.48(8H,m), 7.55-7.57(1H,m), 7.61(2H,d,J=8.6Hz), 9.09(1H,brs), 9.31(1H,d,J=1.5Hz), 9.65(1H,brs), 10.79(1H,s)

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Table 2 (Continuation)

20	mp: 140-143°C. NMR δ : 3.01-3.09(3H,m), 3.16(3H,brs), 3.93(2H,s), 5.06(1H,d,J=10.3Hz), 5.47(2H,s), 6.15(1H,brs), 7.19(2H,d,J=8.6Hz), 7.29-7.33(1H,m), 7.38-7.46(7H,m), 7.61(2H,d,J=8.6Hz), 7.63(1H,s), 7.70(1H,s), 9.08(1H,brs), 9.38(1H,s), 9.63(1H,brs), 10.78(1H,s)
21	mp: 141-146°C. NMR δ : 2.96-3.14(3H,m), 3.15(3H,brs), 3.91(2H,s), 5.04(1H,d,J=10.3Hz), 5.45(2H,s), 6.22(1H,brs), 7.19(2H,d,J=8.6Hz), 7.29-7.42(6H,m), 7.50(3H,s), 7.59(2H,d,J=8.6Hz), 7.65(1H,s), 9.02(1H,brs), 9.32(1H,d,J=1.5Hz), 9.55(1H,brs), 10.73(1H,s)
22	mp: 230-235°C. NMR δ : 2.59-3.10(3H,m), 3.10-3.25(3H,m), 4.47(2H,s), 5.01(1H,dd,J=10.3, 2.4Hz), 5.45(2H,s), 6.21(1H,brs), 7.16-7.22(4H,m), 7.28-7.50(7H,m), 7.54(2H,d,J=8.3Hz), 7.63(2H,dd,J=5.8, 1.9Hz), 8.94(1H,brs), 9.42(1H,brs), 10.98(1H,s)
23	mp: 203-209°C. NMR δ : 2.90-3.10(3H,m), 3.10-3.20(3H,m), 4.41-4.48(2H,m), 4.95-5.05(1H,m), 5.46(2H,s), 6.21(1H,brs), 7.20(2H,d,J=8.6Hz), 7.30-7.42(6H,m), 7.50-7.54(2H,m), 7.70(2H,s), 8.92(1H,brs), 9.39(1H,brs), 10.88-10.95(1H,m)
24	mp: 221-223°C. NMR δ : 2.90-3.08(3H,m), 3.10-3.22(3H,m), 4.04(2H,s), 4.97(1H,d,J=9.1Hz), 5.44(2H,s), 6.20(1H,brs), 7.20(2H,d,J=8.1Hz), 7.30-7.41(9H,m), 7.49(2H,d,J=8.6Hz), 7.55(2H,d,J=8.6Hz), 8.83(1H,brs), 9.16(1H,brs), 10.76(1H,s)
25	mp: 222-225°C. NMR δ : 2.60-3.05(3H,m), 3.10-3.20(3H,m), 4.43(2H,s), 5.01(1H,d,J=7.6Hz), 5.44(2H,s), 6.21(1H,brs), 7.15-7.23(4H,m), 7.26-7.46(5H,m), 7.51(2H,d,J=8.8Hz), 7.65-7.72(4H,m), 8.94(1H,brs), 9.41(1H,brs), 10.93(1H,s), 14.72(1H,brs)
26	mp: 197-203°C. NMR δ : 2.80-3.10(3H,m), 3.10-3.25(3H,m), 4.44(2H,s), 4.99(1H,d,J=8.0Hz), 5.61(2H,s), 6.21(1H,brs), 7.17(2H,d,J=8.6Hz), 7.30-7.42(5H,m), 7.48(2H,d,J=8.5Hz), 7.54(2H,d,J=8.0Hz), 7.70(2H,d,J=8.1Hz), 7.72-7.77(2H,m), 8.90(1H,brs), 9.34(1H,brs), 10.90(1H,s)
27	mp: 208-214°C. NMR δ : 2.90-3.10(3H,m), 3.10-3.22(3H,m), 4.44(2H,s), 4.97(1H,d,J=9.7Hz), 5.62(2H,s), 6.20(1H,brs), 7.16(2H,d,J=8.0Hz), 7.30-7.55(10H,m), 7.70-7.94(6H,m), 8.82(1H,brs), 9.14(1H,brs), 10.76(1H,s)
28	mp: 219-223°C. NMR δ : 2.11(3H,s), 2.92-3.08(3H,m), 3.10-3.20(3H,m), 4.43(2H,s), 5.02(1H,dd,J=10.2, 2.4Hz), 5.51(2H,s), 6.22(1H,brs), 7.14-7.34(7H,m), 7.36-7.42(4H,m), 7.48-7.53(3H,m), 8.95(1H,brs), 9.43(1H,brs), 10.94(1H,s), 14.61(1H,brs)
29	mp: 204-207°C. NMR δ : 2.24(3H,s), 2.80-3.10(3H,m), 3.10-3.50(3H,m), 4.43(2H,s), 5.01(1H,dd,J=10.3, 2.5Hz), 5.39(2H,s), 6.21(1H,brs), 7.17-7.24(2H,m), 7.30-7.42(7H,m), 7.47(2H,dd,J=8.8, 5.4Hz), 7.55(2H,d,J=8.3Hz), 8.94(1H,brs), 9.40(1H,brs), 11.00(1H,s), 14.70(1H,brs)
30	mp: 225-228°C. NMR δ : 2.90-3.07(3H,m), 3.10-3.23(3H,m), 4.28(2H,s), 4.97(1H,d,J=10.3Hz), 5.68(2H,s), 6.20(1H,d,J=3.4Hz), 7.16-7.23(4H,m), 7.30-7.46(7H,m), 7.53(2H,d,J=8.8Hz), 8.82(1H,brs), 9.11(1H,brs), 10.63(1H,s)
31	mp: 232-235°C. NMR δ : 2.90-3.10(3H,m), 3.10-3.25(3H,m), 4.03(2H,s), 4.98(1H,d,J=10.3Hz), 5.97(2H,s), 6.20(1H,brs), 7.19(2H,d,J=8.3Hz), 7.29-7.42(6H,m), 7.55(2H,d,J=8.3Hz), 7.67-7.77(2H,m), 8.87(1H,brs), 9.22(1H,brs), 10.49(1H,s), 14.61(1H,brs)
32	mp: 233-235°C. NMR δ : 2.90-3.10(3H,m), 3.10-3.25(3H,m), 4.01(2H,s), 4.98(1H,d,J=10.3Hz), 5.91(2H,s), 6.19(1H,brs), 7.17-7.48(11H,m), 7.55(2H,d,J=8.3Hz), 8.85(1H,brs), 9.18(1H,brs), 10.47(1H,s)
33	mp: 240-242°C. NMR δ : 2.90-3.10(3H,m), 3.10-3.25(3H,m), 4.32(2H,s), 4.98(1H,d,J=10.3, 3.4Hz), 5.72(2H,s), 6.20(1H,d,J=3.9Hz), 7.20(2H,d,J=8.3Hz), 7.30-7.40(6H,m), 7.51(2H,d,J=8.8Hz), 7.62(1H,d,J=8.3Hz), 7.67(1H,d,J=2.0Hz), 8.86(1H,brs), 9.17(1H,brs), 10.67(1H,s)
34	mp: 221-224°C. NMR δ : 2.90-3.07(3H,m), 3.10-3.20(3H,m), 4.05(2H,s), 5.00(2H,dd,J=2.7, 10.2Hz), 7.21(2H,d,J=8.6Hz), 7.29-7.42(5H,m), 7.58(2H,d,J=8.6Hz), 8.83(1H,s), 8.91(1H,brs), 9.32(1H,brs), 10.62(1H,s)
35	mp: 222-224°C. NMR δ : 2.89-3.07(3H,m), 3.12-3.21(3H,m), 3.84(2H,s), 4.33(2H,s), 4.98(1H,dd,J=2.4, 10.2Hz), 7.20(2H,d,J=8.3Hz), 7.22-7.42(10H,m), 7.58(2H,d,J=8.3Hz), 8.87(1H,brs), 9.22(1H,brs), 10.44(1H,s)
36	mp: 242-245°C. NMR δ : 2.11(3H,s), 2.99-3.06(3H,m), 3.09-3.21(3H,m), 3.68(2H,s), 5.00(1H,dd,J=2.1, 10.2Hz), 6.02(1H,brs), 6.98(1H,s), 7.18(2H,d,J=8.1Hz), 7.28-7.42(5H,m), 7.58(2H,d,J=8.1Hz), 8.89(1H,brs), 9.30(1H,brs), 10.25(1H,s), 12.10(1H,s)
37	mp: 252-256°C. NMR δ : 2.89(3H,s), 2.91-3.07(3H,m), 3.11-3.21(3H,m), 3.65(2H,s), 4.95-5.02(1H,m), 6.20(1H,brs), 6.58(1H,s), 7.20(2H,d,J=8.6Hz), 7.28-7.42(5H,m), 7.57(2H,d,J=8.6Hz), 8.87(1H,brs), 9.24(1H,brs), 10.39(1H,s), 12.56(1H,s)

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Table 2 (Continuation)

38	mp: >230°C(dec). NMR δ : 2.88-3.22(6H,m), 3.73(2H,s), 3.65(2H,s), 5.00(1H,dd,J=2.0, 10.0Hz), 6.20(1H,brs), 7.12(1H,s), 7.18(2H,d,J=8.8Hz), 7.28-7.42(5H,m), 7.59(2H,d,J=8.8Hz), 8.39(4H,brs), 8.91(1H,brs), 9.32(1H,brs), 10.41(1H,s), 12.60(1H,s)
39	mp: 177-181°C. NMR δ : 2.90-3.10(3H,m), 3.10-3.25(3H,m), 3.67(2H,s), 5.00(1H,dd,J=10.0, 2.0Hz), 6.68(1H,s), 6.97(1H,t,J=7.2Hz), 7.19(2H,d,J=8.4Hz), 7.27-7.42(9H,m), 7.59(2H,d,J=8.0Hz), 8.90(1H,brs), 9.29(1H,brs), 10.29(1H,s), 10.54(1H,brs)
40	mp: 237-243°C. NMR δ : 2.90-3.06(3H,m), 3.06-3.20(3H,m), 4.45(2H,s), 5.01(1H,dd,J=7.8, 2.0Hz), 5.70(2H,s), 6.21(1H,brs), 7.14(2H,d,J=8.8Hz), 7.29-7.42(5H,m), 7.46(2H,d,J=8.8Hz), 7.54(2H,d,J=8.8Hz), 7.77(2H,dd,J=14.4, 2.0Hz), 8.13(2H,d,J=8.4Hz), 8.94(1H,brs), 9.41(1H,brs), 10.95(1H,s)
41	mp: 151-159°C. NMR δ : 2.90-3.10(3H,m), 3.10-3.20(3H,m), 3.76(2H,s), 5.02(1H,dd,J=10.2, 2.7Hz), 6.70(1H,s), 7.20(2H,d,J=8.8Hz), 7.25-7.40(5H,m), 7.59(2H,d,J=8.8Hz), 8.96(1H,brs), 9.21(1H,brs), 9.43(1H,brs), 10.58(1H,s)
42	mp: 205-209°C. NMR δ : 2.90-3.08(3H,m), 3.13-3.23(3H,m), 4.92-4.97(1H,m), 6.20(1H,brs), 7.19-7.42(10H,m), 7.71(2H,d,J=8.8Hz), 8.76(1H,brs), 8.92(1H,brs), 9.65(1H,s)
43	NMR δ : 2.20(3H,s), 2.90-3.07(3H,m), 3.10-3.20(3H,m), 3.74(2H,s), 5.00(1H,dd,J=2.5, 10.3Hz), 7.20(2H,d,J=8.8Hz), 7.28-7.42(5H,m), 7.59(2H,d,J=8.8Hz), 8.91(1H,brs), 9.13(1H,brs), 9.33(1H,brs), 10.58(1H,s)
44	NMR δ : 1.48(6H,s), 2.86-3.22(6H,m), 4.90-4.96(1H,m), 6.19(1H,brs), 6.40(1H,brs), 7.17(2H,d,J=8.8Hz), 7.27-7.41(5H,m), 7.56(2H,d,J=8.8Hz), 8.74(1H,brs), 8.90(1H,brs), 9.53(1H,brs)
45	NMR δ : 1.68-2.12(4H,m), 2.43-2.59(2H,m), 2.91-3.07(3H,m), 3.11-3.20(3H,m), 3.76-3.81(1H,m), 5.00(1H,dd,J=2.5, 10.3Hz), 6.20(1H,brs), 7.19(2H,d,J=8.3Hz), 7.27-7.42(5H,m), 7.60(1H,d,J=8.3Hz), 8.90(1H,brs), 9.33(1H,brs), 10.43(1H,s)
46	NMR δ : 2.88-3.24(6H,m), 3.83(2H,s), 4.95-5.04(1H,m), 6.19(1H,brs), 7.16-7.22(2H,m), 7.26-7.45(6H,m), 7.55-7.63(2H,m), 7.87(1H,s), 8.04(1H,d,J=3.6Hz), 8.91(1H,brs), 9.32(1H,brs), 10.42(1H,brs)
47	MS (m/z): 456[(M+H) ⁺]. NMR δ : 2.84-3.19(6H,m), 4.03(2H,s), 4.87-4.97(1H,m), 5.43(2H,s), 6.12(2H,s), 7.20(2H,d,J=8.3Hz), 7.25-7.41(11H,m), 7.53(2H,d,J=8.3Hz), 7.90(1H,s), 10.38(1H,s)
48	NMR δ : 2.88-3.18(6H,m), 3.69(2H,s), 4.87-4.95(1H,m), 5.36(2H,s), 6.15-6.21(1H,m), 7.18(2H,d,J=8.3Hz), 7.27-7.41(11H,m), 7.54(2H,d,J=8.3Hz), 8.57(1H,s), 8.72(1H,brs), 8.82(1H,brs), 10.20(1H,s)
49	NMR δ : 2.88-3.07(3H,m), 3.11-3.21(3H,m), 3.67(2H,s), 4.93-4.99(1H,m), 5.53(2H,s), 6.20(1H,d,J=3.9Hz), 7.00(1H,s), 7.13(2H,d,J=7.3Hz), 7.18(2H,d,J=8.3Hz), 7.24-7.42(8H,m), 7.49(2H,d,J=8.3Hz), 8.82(1H,brs), 9.11(1H,brs), 10.35(1H,s)
50	NMR δ : 1.76-1.87(2H,m), 2.18-2.26(2H,m), 2.80-3.22(8H,m), 4.39-4.47(1H,m), 4.95-5.07(1H,m), 7.15-7.22(2H,m), 7.27-7.43(5H,m), 7.54-7.63(2H,m), 7.74-7.82(1H,m), 8.27(1H,d,J=7.2Hz), 8.67(1H,d,J=4.8Hz), 8.97(1H,brs), 9.47(1H,brs), 10.74(1H,brs)
51	NMR δ : 2.90-3.10(3H,m), 3.10-3.20(3H,m), 4.18(2H,s), 4.96(1H,d,J=8.0Hz), 6.20(1H,brs), 7.18(2H,d,J=8.6Hz), 7.20-7.60(12H,m), 7.84(1H,s), 7.97(1H,s), 8.83(1H,brs), 9.17(1H,brs), 10.55(1H,s)
52	NMR δ : 1.14(6H,d,J=12.9Hz), 2.83(1H,sep,J=12.9Hz), 2.90-3.22(6H,m), 4.38(2H,s), 4.97(1H,d,J=4.1Hz), 5.39(2H,s), 6.20(1H,brs), 7.07-7.42(10H,m), 7.52(2H,d,J=8.8Hz), 7.67(2H,d,J=3.9Hz), 8.84(1H,brs), 9.17(1H,brs), 10.76(1H,s)
53	NMR δ : 1.14(6H,d,J=12.9Hz), 2.83(1H,sep,J=12.9Hz), 2.90-3.22(6H,m), 4.38(2H,s), 4.97(1H,d,J=4.1Hz), 5.39(2H,s), 6.20(1H,brs), 7.07-7.42(10H,m), 7.52(2H,d,J=8.8Hz), 7.67(2H,d,J=3.9Hz), 8.84(1H,brs), 9.17(1H,brs), 10.76(1H,s)
54	NMR δ : 2.95-3.02(3H,m), 3.15(3H,brs), 4.44(2H,s), 5.01(1H,dd,J=10.3, 2.5Hz), 5.58(2H,s), 6.21(1H,brs), 7.19(2H,d,J=8.6Hz), 7.27-7.42(6H,m), 7.51(2H,d,J=8.6Hz), 7.58-7.60(1H,m), 7.69(1H,d,J=2.4Hz), 7.72(1H,d,J=2.0Hz), 7.75(1H,d,J=2.0Hz), 8.96(1H,brs), 9.44(1H,brs), 10.91(1H,s)
55	NMR δ : 2.94-3.04(3H,m), 3.15(3H,brs), 3.94(2H,s), 5.01(1H,d,J=10.3Hz), 5.31(2H,s), 6.21(1H,d,J=3.9Hz), 7.01(1H,s), 7.17-7.41(12H,m), 7.54(2H,d,J=8.3Hz), 8.98(1H,brs), 9.35(1H,brs), 10.55(1H,s)
56	NMR δ : 2.95-3.05(3H,m), 3.15(3H,brs), 4.44(2H,s), 5.01(1H,dd,J=10.3, 2.5Hz), 5.51(2H,s), 6.20(1H,brs), 7.19(3H,d,J=8.6Hz), 7.26-7.42(7H,m), 7.50-7.54(3H,m), 7.58(1H,d,J=2.0Hz), 7.73(1H,d,J=2.0Hz), 8.95(1H,brs), 9.43(1H,brs), 10.98(1H,s)

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Table 2 (Continuation)

57	NMR δ : 2.92-3.05(3H,m), 3.15(3H,brs), 4.43(2H,s), 5.01(1H,dd,J=10.2, 2.6Hz), 5.65(2H,s), 7.20(2H,d,J=8.4Hz), 7.29-7.48(5H,m), 7.50-7.53(3H,m), 7.70(1H,d,J=2.0Hz), 7.78(1H,d,J=2.0Hz), 7.85(1H,dt,J=8.0, 2.0Hz), 8.49(1H,d,J=8.0Hz), 8.94(1H,brs), 9.42(1H,brs), 10.86(1H,s)
58	mp: 150-152°C. NMR δ : 2.88-3.07(3H,m), 3.08(3H,m), 3.95(2H,s), 5.00(1H,dd,J=2.8, 10.0Hz), 6.21(1H,s), 6.82(1H,d,J=7.6Hz), 6.91(1H,d,J=8.0Hz), 7.17-7.23(2H,m), 7.28-7.43(5H,m), 7.55-7.62(2H,m), 7.82-8.04(3H,m), 8.90(1H,brs), 9.31(1H,brs), 10.67(1H,brs), 14.07(1H,brs)
59	NMR δ : 2.90-3.25(6H,m), 4.95-5.04(1H,m), 5.20(1H,s), 6.22(1H,brs), 6.78(1H,s), 7.17-7.24(2H,m), 7.27-7.44(5H,m), 7.67-7.75(2H,m), 8.50-9.10(3H,br), 9.45(1H,br), 10.22(1H,brs)
60	mp: 214-216°C. NMR δ : 2.86-3.24(6H,m), 3.65(2H,s), 4.98(1H,dd,J=2.8, 10.4Hz), 6.18(1H,d,J=6.8Hz), 6.28(1H,d,J=8.8Hz), 7.16-7.22(2H,m), 7.28-7.45(6H,m), 7.53-7.59(2H,s), 8.85(1H,brs), 9.18(1H,brs), 10.36(1H,brs)
61	mp: 180-182°C. NMR δ : 0.87(6H,d,J=6.8Hz), 2.05-2.15(1H,m), 2.59-3.10(3H,m), 3.10-3.20(3H,m), 4.03(2H,d,J=7.8Hz), 4.41(2H,s), 5.01(1H,d,J=8.3Hz), 6.20(1H,brs), 7.21(2H,d,J=8.3Hz), 7.29-7.42(9H,m), 7.60(2H,d,J=8.8Hz), 7.69(1H,d,J=1.9Hz), 7.75(1H,d,J=2.0Hz)
62	mp: 226-228°C. NMR δ : 2.87-3.23(6H,m), 4.45(2H,s), 5.02(1H,dd,J=2.4, 10.0Hz), 5.55(2H,s), 6.21(1H,brs), 7.16-7.46(11H,m), 7.49-7.55(2H,m), 7.66(1H,d,J=2.0Hz), 7.71(1H,d,J=2.0Hz), 8.95(1H,brs), 9.44(1H,brs), 10.93(1H,brs), 14.82(1H,brs)
63	mp: 224-225°C. NMR δ : 2.90-3.05(3H,m), 3.05-3.25(3H,m), 4.46(2H,s), 5.01(1H,d,J=8.0Hz), 5.50(2H,s), 6.21(1H,brs), 7.14-7.50(11H,m), 7.54(2H,d,J=8.8Hz), 7.70-7.73(2H,m), 8.93(1H,brs), 9.39(1H,brs), 10.95(1H,s)
64	mp: 205-208°C. NMR δ : 2.90-3.06(3H,m), 3.10-3.21(3H,m), 4.41(2H,s), 4.99(1H,d,J=8.3Hz), 5.51(2H,s), 6.21(1H,s), 7.06-7.12(1H,m), 7.20(2H,d,J=8.3Hz), 7.28-7.42(6H,m), 7.69(2H,dd,J=2.0, 8.3Hz), 8.87(1H,s), 9.26(1H,s), 10.81(1H,s)
65	mp: 211-216°C. NMR δ : 3.00(3H,brs), 3.15(3H,brs), 4.44(2H,s), 5.05(1H,dd,J=10.2, 1.9Hz), 5.58(2H,s), 6.22(1H,brs), 7.14-7.22(4H,m), 7.29-7.32(1H,m), 7.37-7.42(4H,m), 7.47-7.54(3H,m), 7.65(1H,s), 7.69(1H,d,J=1.9Hz), 9.02(1H,brs), 9.55(1H,brs), 10.97(1H,s)
66	mp: 199-201°C. NMR δ : 2.87-3.23(6H,m), 4.45(2H,s), 4.95-5.04(1H,m), 5.51(2H,s), 6.20(1H,brs), 7.10-7.43(10H,m), 7.49-7.55(2H,m), 7.71(1H,d,J=2.0Hz), 7.74(1H,d,J=2.0Hz), 8.89(1H,brs), 9.30(1H,brs), 10.90(1H,brs), 14.73(1H,brs)
67	mp: 131-135°C. NMR δ : 3.00(3H,brs), 3.16(3H,brs), 4.49(2H,s), 5.04(1H,d,J=10.0Hz), 5.56(2H,s), 6.23(1H,brs), 7.20(2H,d,J=8.2Hz), 7.23-7.34(4H,m), 7.37-7.42(4H,m), 7.53(2H,d,J=8.2Hz), 7.72(2H,s), 9.01(1H,brs), 9.54(1H,brs), 11.00(1H,s)
68	mp: 217-219°C. NMR δ : 2.90-3.05(3H,m), 3.05-3.20(3H,m), 4.46(2H,s), 5.00(1H,d,J=8.0Hz), 5.47(2H,s), 6.21(1H,brs), 7.20(2H,d,J=8.0Hz), 7.25-7.50(7H,m), 7.50-7.60(3H,m), 7.70(1H,d,J=1.9Hz), 7.71(1H,d,J=2.0Hz), 8.91(1H,brs), 9.33(1H,brs), 10.93(1H,s)
69	mp: 213-217°C. NMR δ : 2.90-3.05(3H,m), 3.05-3.20(3H,m), 4.42(2H,s), 5.02(1H,dd,J=10.2, 2.4Hz), 5.62(2H,s), 6.21(1H,brs), 7.20(2H,d,J=8.3Hz), 7.29-7.42(6H,m), 7.49(2H,d,J=8.3Hz), 7.51-7.60(1H,m), 7.68-7.73(2H,m), 8.95(1H,brs), 9.42(1H,brs), 10.89(1H,s)
70	mp: 212-213°C. NMR δ : 2.87-3.23(6H,m), 4.47(2H,s), 5.02(1H,dd,J=2.4, 10.0Hz), 5.53(2H,s), 6.21(1H,brs), 7.16-7.23(2H,m), 7.28-7.34(1H,m), 7.36-7.43(4H,m), 7.48-7.55(2H,m), 7.57-7.67(2H,m), 7.69-7.74(2H,m), 8.96(1H,brs), 9.43(1H,brs), 10.95(1H,brs), 14.86(1H,brs)
71	mp: 209-213°C. NMR δ : 2.90-3.05(3H,m), 3.05-3.20(3H,m), 4.47(2H,s), 4.98-5.01(1H,m), 5.49(2H,s), 6.21(1H,brs), 7.21(2H,d,J=8.3Hz), 7.28-7.34(1H,m), 7.36-7.44(6H,m), 7.53(2H,d,J=8.8Hz), 7.71(1H,d,J=1.9Hz), 7.74(1H,d,J=1.9Hz), 8.91(1H,brs), 9.34(1H,brs), 10.97(1H,s)
72	mp: 190-193°C. NMR δ : 2.90-3.08(3H,m), 3.10-3.21(3H,m), 4.38(2H,s), 4.99(1H,dd,J=2.5, 10.2Hz), 5.69(2H,s), 6.20(1H,s), 7.21(2H,d,J=8.8Hz), 7.29-7.42(5H,m), 7.48(2H,d,J=8.3Hz), 7.70(1H,d,J=1.9Hz), 7.77(1H,s), 8.88(1H,s), 9.27(1H,s), 10.84(1H,s)
73	mp: 233-234°C. NMR δ : 2.90-3.23(6H,m), 4.47(2H,s), 5.02(1H,dd,J=2.4, 10.0Hz), 5.44(2H,s), 6.21(1H,brs), 7.12-7.23(3H,m), 7.28-7.34(1H,m), 7.36-7.44(5H,m), 7.52-7.58(2H,m), 7.66-7.73(3H,m), 7.79-7.81(1H,m), 8.96(1H,brs), 9.44(1H,brs), 10.96(1H,brs), 14.79(1H,brs)

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Table 2 (Continuation)

74	mp: 180-183°C. NMR δ : 2.67-2.76(4H,m), 2.78-2.86(2H,m), 4.00(2H,s), 4.66(1H,dd,J=8.3, 3.9Hz), 5.39(2H,s), 5.42(1H,brs), 6.57(1H,d,J=0.9Hz), 6.78(1H,s), 7.03(2H,d,J=8.3Hz), 7.21-7.26(1H,m), 7.27-7.34(4H,m), 7.46-7.50(1H,m), 7.52(2H,d,J=8.3Hz), 7.56(1H,s), 7.58(1H,s), 8.32(1H,s), 10.32(1H,s)
75	mp: 210-215°C. NMR δ : 2.91-3.03(3H,m), 3.15(3H,brs), 4.44(2H,s), 5.01(1H,dd,J=10.4, 2.6Hz), 5.53(2H,s), 6.21(1H,brs), 7.18(2H,d,J=8.3Hz), 7.30-7.32(1H,m), 7.37-7.42(4H,m), 7.48(2H,d,J=8.3Hz), 7.49(2H,d,J=8.3Hz), 7.74(1H,d,J=2.0Hz), 7.75(1H,d,J=2.0Hz), 7.79(2H,d,J=8.3Hz), 8.94(1H,brs), 9.39(1H,brs), 10.93(1H,s)
76	mp: 162-165°C. NMR δ : 2.93-3.05(3H,m), 3.14(3H,brs), 4.47(2H,s), 5.03(1H,dd,J=10.3, 2.5Hz), 5.62(1H,brs), 5.89(2H,s), 7.12(2H,d,J=8.3Hz), 7.30-7.37(1H,m), 7.39-7.43(6H,m), 7.61(2H,d,J=8.8Hz), 7.69(1H,t,J=7.5Hz), 7.75(1H,d,J=1.9Hz), 7.83-7.86(2H,m), 7.97(1H,d,J=8.3Hz), 8.44(1H,d,J=8.3Hz), 8.99(1H,brs), 9.52(1H,brs), 10.84(1H,s)
77	NMR δ : 2.64-2.74(4H,m), 2.77-2.82(2H,m), 3.93(2H,s), 4.63(1H,dd,J=7.8, 4.4Hz), 5.33(2H,s), 6.80(2H,d,J=6.3Hz), 7.14(2H,d,J=8.8Hz), 7.20-7.24(1H,m), 7.28-7.35(5H,m), 7.43(1H,d,J=7.8Hz), 7.47-7.52(3H,m), 10.27(1H,s)
78	NMR δ : 2.63-2.72(4H,m), 2.75-2.81(2H,m), 3.79(2H,s), 4.62(1H,dd,J=7.8, 4.4Hz), 5.30(1H,brs), 5.33(2H,s), 6.68(1H,d,J=1.0Hz), 6.91(1H,dd,J=8.8, 5.9Hz), 7.06(1H,d,J=1.0Hz), 7.12(2H,d,J=8.8Hz), 7.19-7.24(2H,m), 7.28-7.33(4H,m), 7.43(2H,d,J=8.3Hz), 7.49(1H,dd,J=8.3, 2.5Hz), 8.32(1H,s), 10.21(1H,s)
79	NMR δ : 2.88-3.08(3H,m), 3.10-3.22(3H,m), 4.40(2H,s), 4.97(1H,d,J=8.3Hz), 5.56(2H,s), 6.20(1H,s), 7.19(2H,d,J=8.3Hz), 7.24(1H,d,J=2.5Hz), 7.30-7.60(9H,m), 7.64(1H,d,J=2.0Hz), 7.72(1H,s), 8.83(1H,s), 9.14(1H,s), 10.71(1H,s)
80	NMR δ : 2.90-3.08(3H,m), 3.10-3.22(3H,m), 4.44(2H,s), 5.02(1H,d,J=8.8Hz), 5.59(2H,s), 6.21(1H,s), 7.20(2H,d,J=8.0Hz), 7.24-7.42(7H,m), 7.50(2H,d,J=8.8Hz), 7.72(2H,d,J=6.8Hz), 8.94(1H,s), 9.42(1H,s), 10.93(1H,s)
81	NMR δ : 2.87-3.23(6H,m), 3.85(3H,s), 4.30(2H,s), 4.94-5.01(1H,m), 5.55(2H,s), 6.17-6.22(1H,br), 7.14-7.23(2H,m), 7.28-7.50(9H,m), 7.57-7.64(2H,m), 7.87-7.93(2H,m), 8.83(1H,brs), 9.10(1H,brs), 10.68(1H,brs), 14.86(1H,brs)
82	NMR δ : 1.30-1.64(6H,m), 2.88-3.22(8H,m), 3.45-3.65(2H,m), 4.39(2H,s), 4.97(1H,d,J=9.8Hz), 5.50(2H,s), 6.21(1H,s), 7.20(2H,d,J=8.3Hz), 7.30-7.42(9H,m), 7.51(2H,d,J=8.7Hz), 7.71(2H,d,J=7.8Hz), 8.81(1H,s), 9.14(1H,s), 10.77(1H,s)
83	mp: 229-232°C. NMR δ : 2.90-3.00(3H,m), 3.10-3.18(3H,m), 5.00(1H,dd,J=2.8, 10.1Hz), 5.03(2H,s), 6.27(1H,t,J=2.0Hz), 7.20(2H,d,J=8.8Hz), 7.29-7.42(5H,m), 7.46(1H,d,J=2.4Hz), 7.58(2H,d,J=8.8Hz), 7.77(1H,d,J=2.0Hz), 8.91(1H,s), 9.32(1H,s), 10.53(1H,s)
84	mp: 237-240°C. NMR δ : 2.90-3.08(3H,m), 3.10-3.22(3H,m), 4.96(1H,dd,J=2.0, 10.0Hz), 5.15(2H,s), 7.21(2H,d,J=8.0Hz), 7.28-7.42(5H,m), 7.56(2H,d,J=8.4Hz), 8.03(1H,s), 8.61(1H,s), 8.82(1H,s), 9.09(1H,s), 10.57(1H,s)
85	mp: 244-248°C. NMR δ : 2.90-3.06(3H,m), 3.10-3.20(3H,m), 5.00(1H,d,J=7.6Hz), 5.20(2H,s), 6.20(1H,s), 7.20-7.50(11H,m), 7.59(2H,d,J=7.2Hz), 8.94(3H,s), 9.36(1H,s), 10.95(1H,s), 12.92(1H,s)
86	mp: 223-224°C. NMR δ : 2.86-3.22(6H,m), 3.49(2H,s), 4.93-5.03(1H,m), 6.20(1H,d,J=4.0Hz), 7.15-7.43(9H,m), 7.55-7.62(2H,m), 7.75(1H,dt,J=1.6, 8.0Hz), 8.45-8.53(1H,m), 8.06-9.50(2H,br), 10.35(1H,brs)
87	mp: 236-238°C. NMR δ : 2.86-3.23(6H,m), 3.72(2H,s), 4.91-5.02(1H,m), 6.20(1H,d,J=4.0Hz), 7.15-7.22(2H,m), 7.27-7.45(6H,m), 7.53-7.62(2H,m), 7.73-7.82(1H,m), 8.40-8.60(2H,m), 8.84(1H,brs), 9.16(1H,brs), 10.35-10.50(1H,br)
88	mp: 195-198°C. NMR δ : 2.86-3.22(6H,m), 3.73(2H,s), 4.93-5.04(1H,m), 6.15-6.25(1H,br), 7.14-7.22(2H,m), 7.28-7.43(7H,m), 7.54-7.63(2H,m), 8.47-8.53(2H,m), 9.07(2H,brs), 10.50(1H,brs)
89	mp: 202-204°C. NMR δ : 2.71-2.81(2H,m), 2.88-3.24(8H,m), 3.49(2H,s), 4.93-5.05(1H,m), 6.20(1H,brd,J=3.2Hz), 7.15-7.23(3H,m), 7.26-7.44(6H,m), 7.52-7.60(2H,m), 7.69(1H,dt,J=1.6, 7.6Hz), 8.45-8.51(1H,m), 9.07(2H,brs), 10.07(1H,brs)
90	mp: 220-227°C. NMR δ : 2.80-3.20(8H,m), 4.31(2H,s), 4.42(2H,t,J=8.0Hz), 5.00(1H,d,J=1.0Hz), 6.21(1H,brs), 7.20-7.40(12H,m), 7.59(2H,d,J=8.6Hz), 7.65(2H,dd,J=12.9, 0.9Hz), 8.91(1H,brs), 9.34(1H,brs), 10.98(1H,s)
91	mp: 158-165°C. NMR δ : 2.51-2.78(6H,m), 3.96(2H,s), 4.59(1H,t,J=5.2Hz), 5.20(1H,brs), 7.13-7.32(9H,m), 7.50-7.53(4H,m), 10.33(1H,s), 12.37(1H,brs)
92	mp: 216-217°C. NMR δ : 2.31(3H,s), 2.86-3.24(6H,m), 3.89(2H,s), 4.92-5.07(1H,m), 6.20(1H,d,J=4.0Hz), 7.12-7.22(3H,m), 7.28-7.45(5H,m), 7.50-7.64(2H,m), 8.30(1H,d,J=4.4Hz), 8.60-9.50(2H,br), 10.32(1H,brs)

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Table 2 (Continuation)

93	mp: 236-238°C. NMR δ : 2.86-3.24(6H,m), 3.95(2H,s), 4.91-5.01(1H,m), 5.44(2H,s), 6.19(1H,d,J=4.4Hz), 7.15-7.22(2H,m), 7.27-7.43(5H,m), 7.52-7.62(2H,m), 8.50-8.69(3H,m), 8.83(1H,br), 9.12(1H,brs), 10.41(1H,brs)
94	NMR δ : 2.90-3.10(3H,m), 3.10-3.20(3H,m), 4.38(2H,s), 4.98(1H,t,J=10.4Hz), 5.44(2H,s), 6.20(1H,d,J=3.2Hz), 7.20(2H,d,J=8.4Hz), 7.30-7.45(9H,m), 7.53(2H,d,J=8.8Hz), 7.64(2H,s), 8.85(1H,brs), 9.21(1H,brs), 10.79(1H,s)
95	NMR δ : 2.31(3H,s), 2.89-3.17(6H,m), 3.79(2H,s), 4.98(1H,dt,J=3.2, 10.4Hz), 7.10-7.41(12H,m), 10.32(1H,s)
96	NMR δ : 2.27(3H,s), 2.89-3.17(6H,m), 3.79(2H,s), 4.99(1H,dt,J=3.6, 10.0Hz), 7.17-7.59(12H,m), 10.31(1H,s)
97	NMR δ : 2.44(3H,s), 2.78-3.20(6H,m), 3.80(2H,s), 4.97(1H,dt,J=3.2, 10.4Hz), 7.12-7.66(12H,m), 10.33(1H,s)
98	NMR δ : 1.06(3H,d,J=6.4Hz), 2.50-2.65(2H,m), 2.90-3.15(3H,m), 3.83(2H,s), 4.80-4.94(1H,m), 7.10-7.18(2H,m), 7.23-7.45(7H,m), 7.52-7.60(2H,m), 7.71-7.80(1H,m), 8.41-8.52(1H,m), 10.25(1H,brs)
99	mp: 203-204°C. NMR δ : 1.13(3H,d,J=6.4Hz), 2.55-2.64(1H,m), 3.00-3.50(4H,m), 3.84(2H,s), 4.92-5.02(1H,m), 6.20(1H,d,J=4.0Hz), 7.13-7.20(2H,m), 7.24-7.46(7H,m), 7.54-7.60(2H,m), 7.73-7.80(1H,m), 8.51(1H,brs), 8.67(1H,brs), 9.13(1H,brs), 10.31(1H,brs)
100	NMR δ : 1.06(3H,d,J=6.4Hz), 2.50-2.65(1H,m), 2.57-3.50(4H,m), 3.78(2H,s), 4.77-4.92(1H,m), 5.25(2H,s), 6.85(1H,s), 7.10-7.55(15H,m), 10.33(1H,brs)
101	mp: 194-196°C. NMR δ : 2.88-3.25(6H,m), 3.89(2H,s), 5.20-5.26(1H,m), 6.30(1H,s), 7.17-7.48(7H,m), 7.54-7.60(3H,m), 7.81-7.88(1H,m), 8.54(1H,d,J=4.0Hz), 8.82(1H,s), 9.16(1H,s), 10.35(1H,s)
102	mp: 214-215°C. NMR δ : 2.88-3.25(6H,m), 3.85(2H,s), 4.96-5.02(1H,m), 6.33(1H,d,J=3.8Hz), 7.12-7.31(6H,m), 7.39-7.48(2H,m), 7.58(2H,d,J=8.3Hz), 7.74-7.80(1H,m), 8.50(1H,s), 8.82(1H,s), 9.01(1H,s), 10.30(1H,s)
103	mp: 223-225°C. NMR δ : 2.88-3.06(3H,m), 3.10-3.20(3H,m), 3.84(2H,s), 4.94-5.01(1H,m), 6.24(1H,d,J=4.0Hz), 7.16-7.30(5H,m), 7.38-7.46(3H,m), 7.58(2H,d,J=8.8Hz), 7.76(1H,dt,J=1.6, 7.6Hz), 8.50(1H,d,J=8.8Hz), 8.83(1H,s), 9.08(1H,s), 10.31(1H,s)
104	mp: 208-210°C. NMR δ : 2.88-3.24(6H,m), 3.99(2H,s), 4.90-5.01(1H,m), 6.20(1H,d,J=3.6Hz), 7.15-7.24(2H,m), 7.28-7.44(6H,m), 7.53-7.62(2H,m), 8.50-9.30(4H,m), 10.33(1H,brs)
105	mp: 234-235°C. NMR δ : 2.94-3.25(6H,m), 4.07(2H,s), 4.90-5.02(1H,m), 6.20(1H,d,J=4.0Hz), 7.16-7.23(2H,m), 7.27-7.44(5H,m), 7.53-7.65(4H,m), 7.71-7.78(1H,m), 7.94-8.00(2H,m), 8.33(1H,d,J=8.0Hz), 8.50-9.25(2H,m), 10.46(1H,brs)
106	mp: 221-222°C. NMR δ : 2.90-3.25(6H,m), 3.85(2H,s), 4.92-5.08(1H,m), 6.35(1H,d,J=3.6Hz), 7.14-7.23(2H,m), 7.23-7.31(1H,m), 7.33-7.50(5H,m), 7.54-7.64(2H,m), 7.76(1H,dt,J=1.6, 7.6Hz), 8.43-8.55(1H,m), 8.80-9.40(2H,br), 10.36(1H,brs)
107	mp: 204-205°C. NMR δ : 2.85-3.28(6H,m), 3.85(2H,s), 5.02-5.14(1H,m), 6.37(1H,d,J=4.0Hz), 7.14-7.32(3H,m), 7.36-7.46(2H,m), 7.55-7.64(2H,m), 7.70-7.86(2H,m), 8.46-8.56(2H,m), 8.57-8.65(1H,m), 9.13(2H,brs), 10.37(1H,brs)
108	NMR δ : 2.63-2.67(4H,m), 2.73-2.78(2H,m), 4.07(2H,s), 4.60(1H,dd,J=7.4, 4.9Hz), 5.24(1H,brs), 5.57(2H,s), 7.12-7.23(7H,m), 7.27-7.31(4H,m), 7.37(3H,d,J=8.3Hz), 7.46(2H,d,J=8.3Hz), 7.60-7.61(1H,m), 8.31(1H,s), 10.31(1H,s)
109	NMR δ : 2.26(3H,s), 2.40(3H,s), 2.90-3.17(6H,m), 3.75(2H,s), 4.99(1H,dt,J=3.2, 6.8Hz), 6.97-7.60(11H,m), 10.35(1H,s)
110	mp: 183-184°C. NMR δ : 1.85-2.05(2H,m), 2.53-2.65(2H,m), 2.83-3.03(3H,m), 3.05-3.16(1H,m), 3.88(2H,s), 4.95(1H,d,J=9.6Hz), 6.15(1H,brs), 7.10-7.18(2H,m), 7.22-7.43(7H,m), 7.50-7.60(2H,m), 7.75(1H,dt,J=1.6, 7.2Hz), 8.45-8.53(1H,m), 8.91(2H,brs), 10.29(1H,brs)
111	mp: 225-226°C. NMR δ : 3.02-3.14(1H,m), 3.18-3.46(3H,m), 3.84(2H,s), 4.22-4.35(2H,m), 4.98-5.08(1H,m), 6.21(1H,d,J=3.6Hz), 6.90-6.97(2H,m), 7.23-7.44(7H,m), 7.53-7.62(2H,m), 7.76(1H,dt,J=1.6, 7.2Hz), 8.45-8.54(1H,m), 8.80-9.50(2H,br), 10.29(1H,brs)
112	NMR δ : 1.21(6H,s), 2.85-3.23(4H,m), 3.89(2H,s), 4.90-5.00(1H,m), 6.21(1H,brs), 7.11-7.19(2H,m), 7.28-7.50(7H,m), 7.53-7.62(2H,m), 7.78-7.90(1H,m), 8.45-8.60(2H,m), 9.00-9.10(1H,br), 10.35(1H,brs)
113	mp: 132-133°C. NMR δ : 2.90-3.10(3H,m), 3.13-3.23(3H,m), 4.96(1H,dd,J=2.5, 10.2Hz), 7.06-7.11(1H,m), 7.21(2H,d,J=8.7Hz), 7.30-7.42(5H,m), 7.47-7.53(3H,m), 7.81-7.87(1H,m), 8.29(1H,d,J=4.9Hz), 8.78(1H,s), 9.00(1H,s), 9.88(1H,s), 10.51(1H,s)

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Table 3

Ex.	Structure
1	
2 3	
3 3	
4 1	
4 7	
5 8	
8 6	
9 3	
1 0 4	

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The compounds shown in Tables 4 and 5 together with chemical structural formulae can be easily manufactured by almost the same method as mentioned in the above Examples or Manufacturing Methods or by the method to which some modifications known to the persons skilled in the art are applied. Incidentally, in some cases, there are tautomeric, geometric or optical isomers for the compounds mentioned in Tables 4 and 5, and the compounds of the present invention cover each of the isolated isomers of the above-mentioned ones or a mixture thereof.

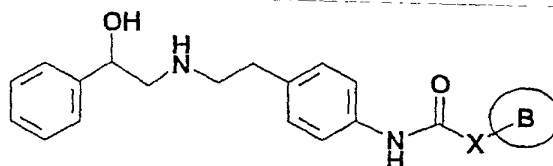
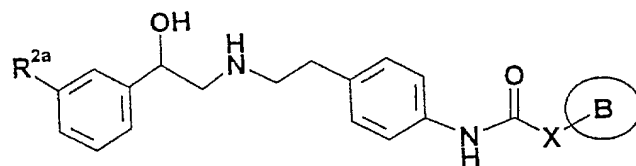


Table 4

No.		No.		No.	
1		2		3	
4		5		6	
7		8		9	
10		11		12	

Table 5

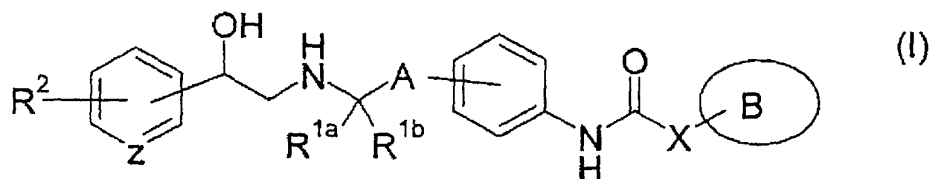


No.	R ^{2a}	X-B	No.	R ^{2a}	X-B
13	H		14	H	
15	H		16	H	
17	H		18	H	
19	H		20	H	
21	Cl		22	Cl	

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Claims

1. An amide derivative represented by the following general formula (I):



(in the formula, each of the symbols means as follows:

ring B: a heteroaryl group which may be substituted and may be fused with a benzene ring;

X: a bond, lower alkylene or alkenylene which may be substituted with hydroxy or a lower alkyl group, carbonyl, or a group represented by -NH- (when X is a lower alkylene group which may be substituted with a lower alkyl group, the hydrogen atoms bonded to the carbon atom constituting the ring B may form a lower alkylene group together with the lower alkyl group so that a ring is formed);

A: lower alkylene or a group represented by -lower alkylene-O-;

R^{1a}, R^{1b}: they may be the same or different and each is a hydrogen atom or a lower alkyl group;

R²: a hydrogen atom or a halogen atom; and

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Z: a nitrogen atom or a group represented by =CH-)

or a salt thereof.

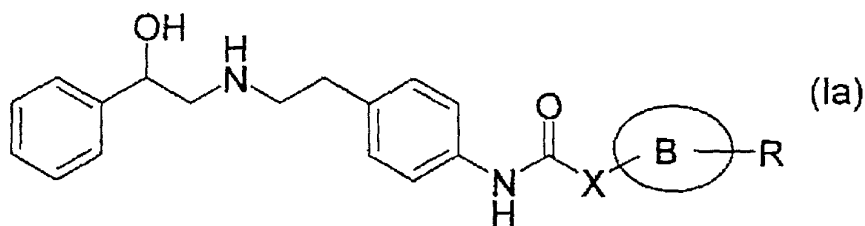
2. The amide derivative or the salt thereof according to claim 1, wherein A is methylene, ethylene, or a group represented by -CH₂O-.

3. The amide derivative or the salt thereof according to claim 2, wherein the ring B is a heteroaryl group which may be substituted with a substituent selected from a halogen atom, lower alkyl, lower alkenyl, lower alkynyl, hydroxy, sulfanyl, halogeno lower alkyl, lower alkyl-O-, lower alkyl-S-, lower alkyl-O-CO-, carboxy, sulfonyl, sulfinyl, lower alkyl-SO-, lower alkyl-SO₂-, lower alkyl-CO-, lower alkyl-CO-O-, carbamoyl, lower alkyl-NH-CO-, di-lower alkyl-N-CO-, nitro, cyano, amino, lower alkyl-NH-, di-lower alkyl-N-, aryl-lower alkyl, halogeno aryl-lower alkyl, guanidino, lower alkyl-CO-NH, and lower alkyl-SO₂-NH-.

4. The amide derivative or the salt thereof according to claim 3, wherein R², R^{1a} and R^{1b} are each a hydrogen atom, and Z is =CH-.

5. An amide derivative represented by the following general formula (Ia):

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(in the formula, each of the symbols means as follows:

ring B: a heteroaryl group;

X: a bond or a lower alkylene group

R: a hydrogen atom, a halogen atom, a lower alkyl group, amino group, an aryl lower alkyl group, or a halogeno aryl-lower alkyl group)

or a salt thereof.

6. (R)-4'-[2-[(2-Hydroxy-2-phenylethyl)amino]ethyl]-2-pyridinecarboxyanilide, (R)-2-[1-(4-chlorobenzyl)-1H-imidazol-2-yl]-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]-acetanilide, (R)-2-[1-(3,4-dichlorobenzyl)-1H-tetrazol-5-yl]-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide, (R)-2-(2-aminothiazol-4-yl)-4'-[2-(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide, (R)-2-(2-benzyl-1H-1,2,4-triazol-3-yl)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide, (R)-2-(2-aminopyridin-6-yl)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]acetanilide, (R)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]-2-(2-pyridyl)acetanilide, (R)-4'-[2-[(2-hydroxy-2-phenylethyl)amino]ethyl]-2-(2-pyrazinyl)acetanilide, (R)-4'-[2-[(2-

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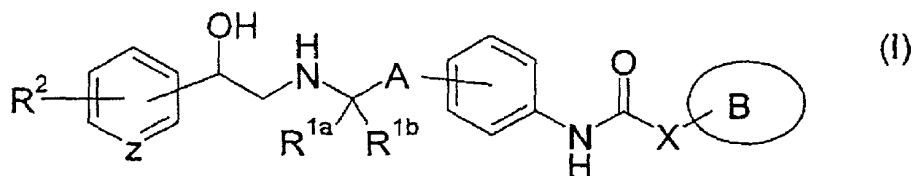
hydroxy-2-phenylethyl)amino]ethyl]-2-(2-pyrimidinyl)-
acetanilide, and salts thereof.

7. A pharmaceutical agent comprising the amide derivative
or the salt thereof according to claims 1 through 6.

8. A therapeutic agent for diabetes mellitus comprising the
amide derivative or the salt thereof according to claims 1
through 6 as an effective ingredient.

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Abstract



Amide derivatives represented by general formula (I) or salts thereof wherein each symbol has the following meaning: ring B: an optionally substituted heteroaryl optionally fused with a benzene ring; X: a bond, lower alkylene or lower alkenylene optionally substituted by hydroxy or lower alkyl, carbonyl, or a group represented by -NH- (when X is lower alkylene optionally substituted by lower alkyl which may be bonded to the hydrogen atom bonded to a constituent carbon atom of ring B to form lower alkylene to thereby form a ring); A: a lower alkylene or a group represented by -(lower alkylene)-O-; R^{1a} and R^{1b}: the same or different and each hydrogen or lower alkyl; R²: hydrogen or halogeno; and Z: nitrogen or a group represented by =CH-. The compounds are useful as a diabetes remedy which not only functions to both accelerate the secretion of insulin and enhance insulin sensitivity but has an antiobestic action and an antihyperlipemic action based on its selective stimulative action on a β_3 receptor.

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Declaration and Power of Attorney for Patent Application

特許出願宣言書

Japanese Language Declaration

私は、下欄に氏名を記載した発明者として、以下のとおり宣言する：

私の住所、郵便の宛先及び国籍は、下欄に氏名に続いて記載したとおりであり、

名称の発明に関し、請求の範囲に記載した特許を求める主題の本来の、最初にして唯一の発明者である（一人の氏名のみが下欄に記載されている場合）か、もしくは本来の、最初にして共同の発明者である（複数の氏名が下欄に記載されている場合）と信じ、

アミド誘導体 又はその塩

その明細書を
(該当する方に印を付す)

☐ ここに添付する。

☒ 1998年10月15 日に出願番号
第 PCT/JP98/04671 号として提出し、
日に補正した。
(該当する場合)

私は、前記のいかなる補正書により補正した前記明細書の内容を検討し、理解したことを陳述する。

私は、連邦規則法典第37部第1章第56条(a)項に従い、本願の審査に所要の情報を開示すべき義務を有することを認める。

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled.

AMIDE DERIVATIVES OR SALTS THEREOF

the specification of which

(check one)

☐ is attached hereto.

☒ was filed on October 15, 1998 as
Application Serial No. PCT/JP98/04671
and was amended on
(if applicable)

I hereby state that I have reviewed and understand the contents of the above identified specification, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the examination of this application in accordance with Title 37, Code of Federal Regulations, §1.56(a).

Japanese Language Declaration

私は、合衆国法典第35部第119条にもとづく下記のいかなる外国特許出願または発明者証出願の外国優先権利益を主張し、さらに優先権の主張に係わる出願の出願日前の出願日を有するいかなる外国特許出願または発明者証出願を以下に明記する：

Prior foreign applications
先の外国出願

Pat. Hei-9-285778	Japan	17, October, 1997 ✓
(Number) (番号)	(Country) (国名)	(Day / Month / Year Filed) (出願の日/月/年)
(Number) (番号)	(Country) (国名)	(Day / Month / Year Filed) (出願の日/月/年)
(Number) (番号)	(Country) (国名)	(Day / Month / Year Filed) (出願の日/月/年)

Priority claimed
優先権の主張

<input checked="" type="checkbox"/> Yes あり	<input type="checkbox"/> No なし
<input type="checkbox"/> Yes あり	<input type="checkbox"/> No なし
<input type="checkbox"/> Yes あり	<input type="checkbox"/> No なし

私は、合衆国法典第35部第120条にもとづく下記の合衆国特許出願の利益を主張し、本願の請求の範囲各項に記載の主題が合衆国法典第35部第112条第1項に規定の態様で先の合衆国出願に開示されていない限度において、先の出願の出願日と本願の国内出願日またはPCT国際出願日の間に公表された連邦規則法典第37部第1章第56条(a)項に記載の所要の情報を開示すべき義務を有することを認める：

I hereby claim the benefit under Title 35, United States Code, §120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, §112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, §1.56(a) which occurred between the filing date of the prior application and the national or PCT international filing date of this application:

(Application Serial No.) (出願番号)	(Filing Date) (出願日)	(現況) (特許済み、係属中、放棄済み)	(Status) (Patented, pending, abandoned)
(Application Serial No.) (出願番号)	(Filing Date) (出願日)	(現況) (特許済み、係属中、放棄済み)	(Status) (Patented, pending, abandoned)

私は、ここに自己の知識にもとづいて行った陳述がすべて真実であり、自己の有する情報及び信ずるところに従って行った陳述が真実であると信じ、さらに故意に虚偽の陳述等を行った場合、合衆国法典第18部第1001条により、罰金もしくは禁錮に処せられるか、またはこれらの刑が併科され、またかかる故意による虚偽の陳述が本願ないし本願に対して付与される特許の有効性を危くすることがあることを認識して、以上の陳述を行ったことを宣言する。

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application, or any patent issued thereon.

Japanese Language Declaration

委任状：私は、下記発明者として、以下の弁理士及び代理人、またはそのいずれかをここに選任し、本願の手続きを遂行すること並びにこれに関する一切の行為を米国特許商標庁に対して行うことを委任する。(代理人氏名及び登録番号を明記のこと)

POWER OF ATTORNEY: As a named inventor, I hereby appoint the following attorney(s) and/or agent(s) to prosecute this application and transact all business in the Patent and Trademark Office connected therewith. (list agent's name(s) and registration number(s))

FINNEGAN, HENDERSON, FARABOW, GARRETT & DUNNER, Reg. No. 22,540; Douglas B. Henderson, Reg. No. 20,291; Ford F. Farabow, Jr., Reg. No. 20,630; Arthur S. Garrett, Reg. No. 20,338; Donald R. Dunner, Reg. No. 19,023; Brian G. Brunsvold, Reg. No. 22,593; Tipton D. Jennings, IV, Reg. No. 20,645; Jerry D. Voight, Reg. No. 23,020; Laurence R. Hefter, Reg. No. 20,827; Kenneth E. Payne, Reg. No. 23,098; Herbert H. Mintz, Reg. No. 26,691; C. Larry O'Rourke, Reg. No. 26,014; Albert J. Santorelli, Reg. No. 22,610; Michael C. Eimer, Reg. No. 25,857; Richard H. Smith, Reg. No. 20,609; Stephen L. Peterson, Reg. No. 26,325; John M. Romary, Reg. No. 26,331; Bruce C. Zotter, Reg. No. 27,680; Dennis P. O'Reilly, Reg. No. 27,932; Allen M. Sokal, Reg. No. 26,695; Robert D. Bajelsky, Reg. No. 25,387; Richard L. Stroup, Reg. No. 28,478; David W. Hill, Reg. No. 28,220; Thomas L. Irving, Reg. No. 28,619; Charles E. Lipsey, Reg. No. 28,165; Thomas W. Winland, Reg. No. 27,605; Basil J. Lewis, Reg. No. 28,818; Martin I. Fuchs, Reg. No. 28,508; E. Robert Yoches, Reg. No. 30,120; Barry W. Graham, Reg. No. 29,924; Susan Haberman Griffen, Reg. No. 30,907; Richard B. Racine, Reg. No. 30,415; Thomas H. Jenkins, Reg. No. 30,857; Robert E. Converse, Jr., Reg. No. 27,432; Clair X. Mullen, Jr., Reg. No. 20,348; Christopher P. Foley, Reg. No. 31,354; John C. Paul, Reg. No. 30,413; Roger D. Taylor, Reg. No. 28,992; David M. Kelly, Reg. No. 30,953; Kenneth J. Meyers, Reg. No. 25,146; Carol P. Einaudi, Reg. No. 32,220; Walter Y. Boyd, Jr., Reg. No. 31,738; Steven M. Anzalone, Reg. No. 32,095; Jean B. Fordis, Reg. No. 32,984; Barbara C. McCurdy, Reg. No. 32,120; James K. Hammond, Reg. No. 31,964; Richard V. Burgujian, Reg. No. 31,744; J. Michael Jakes, Reg. No. 32,824; Thomas W. Banks, Reg. No. 32,719; M. Paul Barker, Reg. No. 32,013; Bryan C. Diner, Reg. No. 32,409; Christopher P. Isaac, Reg. No. 32,616; Andrew C. Sonu, Reg. No. 33,457; Dirk D. Thomas, Reg. No. 32,600

書類の送付先：

Send Correspondence to:

FINNEGAN, HENDERSON, FARABOW, GARRETT & DUNNER, L.L.P.
1300 I STREET, N.W.
WASHINGTON, D.C. 20005-3315

直通電話連絡先：(名称及び電話番号)

Direct Telephone Calls to: (name and telephone number)

ARTHUR S. GARRETT
202-408-4091

唯一のまたは第一の発明者の氏名 丸山 龍也	Full name of sole or first inventor Tatsuya MARUYAMA
同発明者の署名 丸山 龍也 日付 1999年11月10日	Inventor's signature Date
住所 〒305-8585 日本国茨城県つくば市御幸が丘21 山之内製薬株式会社内	Residence c/o Yamanouchi Pharmaceutical Co., Ltd., 21, Miyukigaoka, Tsukuba-shi,
国籍 日本国	Citizenship IBARAKI 305-8585 JAPAN JPK JAPAN
郵便の宛先 〒305-8585 日本国茨城県つくば市御幸が丘21 山之内製薬株式会社内	Post Office Address c/o Yamanouchi Pharmaceutical Co., Ltd., 21, Miyukigaoka, Tsukuba-shi, IBARAKI 305-8585 JAPAN

第二の共同発明者の氏名(該当する場合) 鈴木 貴之	Full name of second joint inventor, if any Takayuki SUZUKI
同第二発明者の署名 鈴木 貴之 日付 1999年11月10日	Second inventor's signature Date
住所 〒305-8585 日本国茨城県つくば市御幸が丘21 山之内製薬株式会社内	Residence c/o Yamanouchi Pharmaceutical Co., Ltd., 21, Miyukigaoka, Tsukuba-shi,
国籍 日本国	Citizenship IBARAKI 305-8585 JAPAN JPK JAPAN
郵便の宛先 〒305-8585 日本国茨城県つくば市御幸が丘21 山之内製薬株式会社内	Post Office Address c/o Yamanouchi Pharmaceutical Co., Ltd., 21, Miyukigaoka, Tsukuba-shi, IBARAKI 305-8585 JAPAN

(第三またはそれ以降の共同発明者に対しても同様な情報及び署名を提供すること。)

(Supply similar information and signature for third and subsequent joint inventors.)

3-00

第三の共同発明者の氏名 (該当する場合) 恩田 健一	Full name of third joint inventor, if any <u>Kenichi ONDA</u>
同第三発明者の署名 恩田 健一 日付 1999年11月10日	Third inventor's signature Date
住所 〒305-8585 日本国茨城県つくば市御幸ヶ丘21 山之内製薬株式会社内	Residence c/o Yamanouchi Pharmaceutical Co., Ltd., 21, Miyukigaoka, Tsukuba-shi,
国籍 日本国	Citizenship IBARAKI 305-8585 JAPAN JFX JAPAN
郵便の宛先 〒305-8585 日本国茨城県つくば市御幸ヶ丘21 山之内製薬株式会社内	Post Office Address c/o Yamanouchi Pharmaceutical Co., Ltd., 21, Miyukigaoka, Tsukuba-shi, IBARAKI 305-8585 JAPAN

4-00

第四の共同発明者の氏名 (該当する場合) 早川 昌彦	Full name of fourth joint inventor, if any <u>Masahiko HAYAKAWA</u>
同第四発明者の署名 早川 昌彦 日付 1999年11月10日	Fourth inventor's signature Date
住所 〒305-8585 日本国茨城県つくば市御幸ヶ丘21 山之内製薬株式会社内	Residence c/o Yamanouchi Pharmaceutical Co., Ltd., 21, Miyukigaoka, Tsukuba-shi,
国籍 日本国	Citizenship IBARAKI 305-8585 JAPAN JFX JAPAN
郵便の宛先 〒305-8585 日本国茨城県つくば市御幸ヶ丘21 山之内製薬株式会社内	Post Office Address c/o Yamanouchi Pharmaceutical Co., Ltd., 21, Miyukigaoka, Tsukuba-shi, IBARAKI 305-8585 JAPAN

5-00

第五の共同発明者の氏名 (該当する場合) 森友 博幸	Full name of fifth joint inventor, if any <u>Hiroyuki MORITOMO</u>
同第五発明者の署名 森友 博幸 日付 1999年11月10日	Fifth inventor's signature Date
住所 〒305-8585 日本国茨城県つくば市御幸ヶ丘21 山之内製薬株式会社内	Residence c/o Yamanouchi Pharmaceutical Co., Ltd., 21, Miyukigaoka, Tsukuba-shi,
国籍 日本国	Citizenship IBARAKI 305-8585 JAPAN JFX JAPAN
郵便の宛先 〒305-8585 日本国茨城県つくば市御幸ヶ丘21 山之内製薬株式会社内	Post Office Address c/o Yamanouchi Pharmaceutical Co., Ltd., 21, Miyukigaoka, Tsukuba-shi, IBARAKI 305-8585 JAPAN

6-00

第六の共同発明者の氏名 (該当する場合) 君塚 哲也	<u>Tetsuya KIMIZUKA</u> Full name of sixth joint inventor, if any
同第六発明者の署名 君塚 哲也 日付 1999年11月10日	Sixth inventor's signature Date
住所 〒305-8585 日本国茨城県つくば市御幸ヶ丘21 山之内製薬株式会社内	c/o Yamanouchi Pharmaceutical Co., Ltd., 21, Miyukigaoka, Tsukuba-shi,
国籍 日本国	Residence IBARAKI 305-8585 JAPAN JFX JAPAN
〒305-8585 郵便の宛先 日本国茨城県つくば市御幸ヶ丘21 山之内製薬株式会社内	Citizenship JAPAN
	Post Office Address c/o Yamanouchi Pharmaceutical Co., Ltd., 21, Miyukigaoka, Tsukuba-shi, IBARAKI 305-8585 JAPAN

(第七またはそれ以降の共同発明者に対しても同様な情報及び署名を提供すること。)

(Supply similar information and signature for seventh and subsequent joint inventors.)

7-00

第七の共同発明者の氏名 (該当する場合) <u>松井哲夫</u>		<u>Tetsuo MATSUI</u>	
Full name of seventh joint inventor, if any			
同第七発明者の署名 <u>松井哲夫</u> 日付 <u>1999年11月10日</u>		Seventh inventor's signature Date	
〒305-8585 住所 <u>日本国茨城県つくば市御幸が丘21</u> <u>山之内製薬株式会社内</u>		c/o Yamanouchi Pharmaceutical Co., Ltd., 21, Miyukigaoka, <u>Tsukuba-shi</u> , IBARAKI Residence 305-8585 JAPAN <u>SPX</u>	
日本国		JAPAN	
国籍		Citizenship	
〒305-8585 郵便の宛先 <u>日本国茨城県つくば市御幸が丘21</u> <u>山之内製薬株式会社内</u>		c/o Yamanouchi Pharmaceutical Co., Ltd., 21, Miyukigaoka, Tsukuba-shi, IBARAKI Post office Address	
		305-8585 JAPAN	

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